

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: March 8, 2005, 16:04:41 ; Search time 39 Seconds  
(without alignments)  
350.328 Million cell updates/sec

Title: US-10-723-083-2  
Perfect score: 765  
Sequence: 1 MHVHHHSSGIEGRMAPARS.....ENLKDFLLVLPDCWEPVOE 142  
Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues  
Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : PIR 79.\*  
1: pir1.\*  
2: pir2.\*  
3: pir3.\*  
4: pir4.\*

Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	673	88.0	144	1 FQHUGM	granulocyte-macrop
2	548	71.6	144	2 JH0469	granulocyte-macrop
3	544	71.1	144	1 A61632	granulocyte-macrop
4	480.5	62.8	143	1 FQBOGM	granulocyte-macrop
5	473	61.8	144	2 A44936	granulocyte-macrop
6	441	57.6	127	2 I46269	granulocyte-macrop
7	367.5	48.0	153	1 FQMSGM	granulocyte-macrop
8	84.5	11.0	2493	2 S72349	nonstructural poly
9	81	10.6	305	2 A56554	transcription fact
10	81	10.6	359	2 A55839	tetrapyrrole methy
11	80.5	10.5	285	2 A42390	nonstructural poly
12	80.5	10.5	2493	2 S26372	homeotic protein B
13	79	10.3	610	2 A57632	hypothetical prote
14	78.5	10.3	329	2 T45972	hypothetical prote
15	78	10.2	496	2 S25091	cruciferin BnC2 -
16	78	10.2	816	2 S05548	gap protein hunchb
17	78	10.2	1019	2 T00117	dve protein - frui
18	77.5	10.1	420	2 T39712	hypothetical prote
19	77.5	10.1	427	2 T42516	hypothetical prote
20	75.5	9.9	292	2 I51171	transcription fact
21	75.5	9.9	942	2 J21229	protein kinase PKN
22	75.5	9.9	1016	1 A46079	protein kinase C (
23	75	9.8	605	1 Q0B529	BKLF1 protein - hu
24	74.5	9.7	384	2 T41302	ferrochelatase pre
25	74.5	9.7	1305	2 T18548	flax rust resistan
26	73.5	9.6	248	2 T02647	probable MADS-box
27	73.5	9.6	1622	2 JE0378	DNA (cytosine-5)-
28	73	9.5	234	2 T26429	hypothetical prote
29	73	9.5	461	2 S34472	MPH-1 protein - mo

30 73 9.5 943 2 T03306  
31 73 9.5 1183 2 A89135  
32 73 9.5 1496 2 T00499  
33 72.5 9.5 289 2 D83709  
34 72.5 9.5 417 2 C32185  
35 72 9.4 311 2 G86383  
36 72 9.4 447 2 A34582  
37 72 9.4 770 2 P0105  
38 71.5 9.3 530 2 D70476  
39 71.5 9.3 1273 2 T00338  
40 71 9.3 398 1 TVFVVR  
41 71 9.3 611 2 F82951  
42 70.5 9.2 265 2 T46089  
43 70.5 9.2 601 1 B56564  
44 70.5 9.2 633 1 A26030  
45 70.5 9.2 1025 2 H81751

## ALIGNMENTS

## RESULT 1

## FQHUGM

granulocyte-macrophage colony-stimulating factor precursor [validated] - human  
N;Alternate names: colony-stimulating factor 2; GM-CSF  
C;Species: Homo sapiens (man)  
C;Date: 04-Dec-1986 #sequence revision 04-Dec-1986 #text change 09-Jul-2004  
C;Accession: C24636; I59065; A21853; A44175; JCI1090  
R;Miyatake, S.; Otsuka, T.; Yokota, T.; Lee, F.; Arai, K.  
EMBL J. 4, 2561-2568, 1985

A;Title: Structure of the chromosomal gene for granulocyte-macrophage colony stimulating  
A;Reference number: A91015; MUID:86030234; PMID:3876930  
A;Accession: C24636  
A;Molecule type: DNA  
A;Residues: 1-144 <MY>  
A;Cross-references: UNIPROT:P04141; EMBL:X03021; NID:g31858; PIDN:CAA26822.1; PID:g31859  
R;Kauschansk, K.; O'Hara, P.J.; Berkner, K.; Segal, G.M.; Hagen, F.S.; Adamson, J.W.  
Proc. Natl. Acad. Sci. U.S.A. 83, 3101-3105, 1986  
A;Title: Genomic cloning, characterization, and multilineage growth-promoting activity of  
A;Reference number: I59065; MUID:86205844; PMID:3486413  
A;Accession: I59065  
A;Status: translated from GB/EMBL/DDBJ  
A;Molecule type: DNA  
A;Residues: 1-144 <RES>  
A;Cross-references: GB:M13207; NID:g181147; PIDN:AAA98768.1; PID:g181148  
R;Cantrell, M.A.; Anderson, D.; Cerretti, D.P.; Price, V.; McKereghan, K.; Tushinski, R.  
Proc. Natl. Acad. Sci. U.S.A. 82, 6250-6254, 1985  
A;Title: Cloning, sequence, and expression of a human granulocyte/macrophage colony-stim  
A;Reference number: A25169; MUID:85298329; PMID:3898082  
A;Accession: A25169  
A;Molecule type: mRNA  
A;Residues: 1-144 <CAN>  
A;Cross-references: GB:M11734; NID:g181149; PIDN:AAAS2122.1; PID:g181150  
R;Lee, F.; Yokota, T.; Otsuka, T.; Gemmell, L.; Larson, N.; Luh, J.; Arai, K.; Rennick, I  
Proc. Natl. Acad. Sci. U.S.A. 82, 4360-4364, 1985  
A;Title: Isolation of cDNA for a human granulocyte-macrophage colony-stimulating factor I  
A;Reference number: A01853; MUID:85242684; PMID:3925454  
A;Accession: A01853  
A;Molecule type: mRNA  
A;Residues: 1-144 <LEE>  
A;Cross-references: GB:M11220; NID:g183363; PIDN:AAAS2578.1; PID:g183364  
R;Wong, G.G.; Witek, J.S.; Temple, P.A.; Wilkens, K.M.; Leary, A.C.; Luxenberg, D.P.; Jon

Science 228, 810-815, 1985  
A;Title: Human GM-CSF: molecular cloning of the complementary DNA and purification of th  
A;Reference number: A44175; MUID:85218749; PMID:3923623  
A;Accession: A44175  
A;Molecule type: mRNA  
A;Residues: 1-116, 'T', 118-144 <WON>  
A;Cross-references: GB:M10663; NID:g181145; PIDN:AAAS2121.1; PID:g181146  
A;Note: parts of this sequence, including the amino end of the mature protein, were conf  
R;Wen, D.Y.; Huang, B.R.; Cai, L.W.; Si, J.Y.  
Acta Biochim. Biophys. Sin. 25, 651-655, 1993



## RESULT 5

A44936  
granulocyte-macrophage colony-stimulating factor precursor - dog  
C/Species: Canis lupus familiaris (dog)  
C/Date: 17-Feb-1994 #sequence\_revision 17-Feb-1994 #text\_change 09-Jul-2004  
C/Accession: A44936  
R;Nash, R.A.; Schuening, F.; Appelbaum, F.; Hammond, W.P.; Boone, T.; Morris, C.F.; Slid  
Blood 78, 930-937, 1991  
A/Title: Molecular cloning and in vivo evaluation of canine granulocyte-macrophage color  
A/Reference number: A44936; MUID:91329842; PMID:1868252  
A/Accession: A44936  
A/Status: preliminary  
A/Molecule type: mRNA  
A/Residues: 1-144 <NAS>  
A/Cross-references: UNIPROT:P48749; GB:S49738; PIDN:AAAB19466.1; PID:G233567  
A/Note: sequence extracted from NCBI backbone (NCBI:49738, NCBI:P:49739)  
C/Superfamily: granulocyte-macrophage colony-stimulating factor  
F:1-17/Domain: signal sequence #status predicted <SIG>

Query Match 61.8%; Score 473; DB 2; Length 144;  
Best Local Similarity 68.5%; Pred. No. 4e-38;  
Matches 87; Conservative 19; Mismatches 21; Indels 0; Gaps 0;  
QY 16 APARSPSPQPEHVNIAQEARLLNLSRDTAAENNETVEVISEMFDLQEPDTCLOTRLE 75  
DB 18 APTRSPTLVTRPSQHVDAIQEALSLLNSNDVTAVNMKAVKVVSEVDFPBGPTCLETRLQ 77  
QY 76 LYKQGLRGSUTLKGPLTMASHYKQHCPTPTSCATOLIITPESKKNLKDFLLVIPP 135  
DB 78 LYKEGGLQSUTSLKNPLTMANHYKQHCPTPTSPCATINPKSFKNLKDFLFIIPP 137

QY 136 CWEPVOE 142  
DB 138 CWKPVK 144

## RESULT 6

I46269  
granulocyte-macrophage colony stimulating factor - rat (fragment)  
C/Species: Rattus norvegicus (Norway rat)  
C/Date: 14-Feb-1997 #sequence\_revision 14-Feb-1997 #text\_change 09-Jul-2004  
C/Accession: I46269  
R;Smith, L.R.; Lundeen, K.A.; Diveley, J.P.; Carlo, D.J.; Brostoff, S.W.  
Immunogenetics 39, 80, 1994  
A/Title: Nucleotide sequence of the Lewis rat granulocyte-macrophage colony stimulating  
A/Reference number: I46269; MUID:94041474; PMID:8225444  
A/Accession: I46269  
A/Status: preliminary; translated from GB/EMBL/DBJ  
A/Molecule type: mRNA  
A/Residues: 1-127 <SMI>  
A/Cross-references: UNIPROT:P48750; EMBL:U00620; NID:G392779; PIDN:AAAL8281.1; PID:G3927  
C/Superfamily: granulocyte-macrophage colony-stimulating factor

Query Match 57.6%; Score 441; DB 2; Length 127;  
Best Local Similarity 63.0%; Pred. No. 4e-35;  
Matches 80; Conservative 19; Mismatches 28; Indels 0; Gaps 0;  
QY 16 APARSPSPQPEHVNIAQEARLLNLSRDTAAENNETVEVISEMFDLQEPDTCLOTRLE 75  
DB 1 APTRSNPVTRPWKHVDIAKEALSLLNDMPALENEKNEVDIISNFSIQRTCVQTRLK 60  
QY 76 LYKQGLRGSUTLKGPLTMASHYKQHCPTPTSCATOLIITPESKKNLKDFLLVIPP 135  
DB 61 LYKQGLRGNLTUNGALTMASHYQNTCPPTPTDCEIETTFEDFKNLKGLFDFIPP 120

QY 136 CWEPVOE 142  
DB 121 CWKPVK 127

## RESULT 7

Query Match 48.0%; Score 367.5; DB 1; Length 153;  
Best Local Similarity 54.3%; Pred. No. 5.5e-28;  
Matches 69; Conservative 23; Mismatches 32; Indels 3; Gaps 1;

## FQMSGM

granulocyte-macrophage colony-stimulating factor precursor - mouse  
N/Alternate names: colony-stimulating factor 2; GM-CSF; integral membrane protein  
C/Species: Mus musculus (house mouse)  
C/Date: 28-Aug-1985 #sequence\_revision 13-Mar-1997 #text\_change 09-Jul-2004  
C/Accession: I48368; I48369; A24636; A24645; A24644; A01854; A21882  
R;Gough, N.M.; Metcalf, D.; Gough, J.; Grail, D.; Dunn, A.R.  
EMBO J. 4, 645-653, 1985  
A/Title: Structure and expression of the mRNA for murine granulocyte-macrophage colony st  
A/Reference number: I48368; MUID:85230531; PMID:3874057  
A/Accession: I48368  
A/Status: preliminary; translated from GB/EMBL/DBJ  
A/Molecule type: mRNA  
A/Residues: 1-153 <RES>  
A/Cross-references: UNIPROT:P01587; EMBL:X02333; NID:G51103; PIDN:CAA26192.1; PID:G51104  
A/Accession: I48369  
A/Status: preliminary; translated from GB/EMBL/DBJ  
A/Molecule type: mRNA  
A/Residues: 13-153 <RE2>  
A/Cross-references: EMBL:X02333; NID:G51103; PIDN:CAA26193.1; PID:G51106  
R;Miyatake, S.; Otsuka, T.; Yokota, T.; Lee, F.; Arai, K.  
EMBO J. 4, 2561-2568, 1985  
A/Title: Structure of the chromosomal gene for granulocyte-macrophage colony stimulating  
A/Reference number: A91015; MUID:86030234; PMID:3876930  
A/Accession: A24636  
A/Molecule type: DNA; mRNA  
A/Residues: 13-150, 'G', 152-153 <MIY>  
A/Cross-references: GB:X03020; NID:G51098; PIDN:CAA26821.1; PID:G51099  
A/Note: the sequence translated from the mRNA differs from that of the DNA in having 151-  
R;Stanley, E.; Metcalf, D.; Sobieszczuk, P.; Gough, N.M.; Dunn, A.R.  
EMBO J. 4, 2369-2373, 1985  
A/Title: The structure and expression of the murine gene encoding granulocyte-macrophage  
A/Reference number: A24645; MUID:86030235; PMID:3876931  
A/Accession: A24645  
A/Molecule type: DNA  
A/Residues: 13-150, 'G', 152-153 <STA>  
A/Cross-references: GB:X03020; NID:G51098; PIDN:CAA26821.1; PID:G51099  
R;DeLamarter, J.F.; Mermod, J.J.; Liang, C.M.; Eliason, J.F.; Thatcher, D.R.  
EMBO J. 4, 2575-2581, 1985  
A/Reference number: A24644; MUID:86030236; PMID:3902470  
A/Accession: A24644  
A/Molecule type: mRNA  
A/Residues: 13-153 <DE>  
A/Cross-references: GB:X03019; NID:G51100; PIDN:CAA26820.1; PID:G736260  
R;Gough, N.M.; Gough, J.; Metcalf, D.; Kelsso, A.; Grail, D.; Nicola, N.A.; Burgess, A.W.  
Nature 309, 763-767, 1984  
A/Title: Molecular cloning of cDNA encoding a murine haematopoietic growth regulator, gra  
A/Reference number: A01854; MUID:84245825; PMID:6610831  
A/Accession: A01854  
A/Molecule type: mRNA  
A/Residues: 36, 'I', 38-150, 'S', 152-153 <GOU>  
A/Cross-references: GB:X05906; NID:G51096; PIDN:CAA29336.1; PID:G51097  
A/Experimental source: lung tissue  
R;Sparrow, L.G.; Metcalf, D.; Hunkapiller, M.W.; Hood, L.E.; Burgess, A.W.  
Proc. Natl. Acad. Sci. U.S.A. 82, 292-296, 1985  
A/Title: Purification and partial amino acid sequence of asialo murine granulocyte-macrop  
A/Reference number: A21882; MUID:85113187; PMID:3871523  
A/Accession: A21882  
A/Molecule type: protein  
A/Residues: 36, 'I', 38-69 <SPA>  
C/Comment: It is unclear if Met-1 or Met-13 is the initiator.  
C/Genetics:  
A/Introns: 62/3; 76/3; 118/3  
C/Superfamily: granulocyte-macrophage colony-stimulating factor  
C/Keywords: cytokine; glycoprotein; growth factor; macrophage; monomer; T-cell  
F:1-29/Domain: signal sequence #status predicted <SIG>  
F:30-153/Product: granulocyte-macrophage colony-stimulating factor #status predicted <MA  
F:95,104/Binding site: carboxydrate (Asn) (covalent) #status predicted

Query Match 48.0%; Score 367.5; DB 1; Length 153;  
Best Local Similarity 54.3%; Pred. No. 5.5e-28;  
Matches 69; Conservative 23; Mismatches 32; Indels 3; Gaps 1;

QY 16 APARSPSPQTPWEHVNAIQEARLLNLSDTAEMNETVEISEMFDLQEPCTCLOTRLE 75  
DB 30 APTRPSPITVTRPKHGVAEIKEA--LNLDDMPVTINEEVEVVSNEFSFKKUTCQVTRUK 86  
QY 76 LYKQGLRGLSLTKLKGPLTMASHYKHQCPPTPETS CATQIIITFESPKENLKDFLLVIPED 135  
DB 87 IFQGLRGNFTKLKALNMTASYYQYCYPTPETDCTQVTVYADFLDSLKFTLTDIPE 146  
QY 136 CWEPVQE 142  
DB 147 CKPQVK 153  
RESULT 8  
S72349  
nonstructural polyprotein - eastern equine encephalomyelitis virus  
N;Alternate names: nonstructural protein NSP1; nonstructural protein NSP2; nonstructural  
C;Species: eastern equine encephalomyelitis virus  
C;Date: 04-May-1998 #sequence\_revision 15-May-1998 #text\_change 09-Jul-2004  
C;Accession: S72349  
R;Weaver, S.C.; Hagenbaugh, A.; Ballew, L.A.; Netesov, S.V.; Volchkov, V.E.; Chang, G.J.  
Virology 197, 375-390, 1993  
A;Title: A comparison of the nucleotide sequences of eastern and western equine encephal  
A;Reference number: S72349; MUID:94025587; PMID:8105605  
A;Accession: S72349  
A;Status: preliminary  
A;Molecule type: genomic RNA  
A;Residues: 1-2493 <WEA>  
A;Cross-references: UNIPROT:Q88789; EMBL:U01034; NID:g939006; PIDN:AA53734.1; PID:g939006  
A;Note: readthrough of the terminator UGA occurs between the codons AAU for residue 1878  
A;Note: the readthrough stopcodon UGA for residue 1879 is translated as X  
C;Superfamily: Semliki Forest virus nonstructural protein  
Query Match 11.0%; Score 84.5; DB 2; Length 2493;  
Best Local Similarity 26.1%; Pred. No. 21;  
Matches 36; Conservative 17; Mismatches 54; Indels 31; Gaps 7;  
QY 17 PARSPSPQTPWEHVNAIQEARLLNLSDTAEMNETVE----- 57  
DB 1678 PARPSP---PCTSTNG--STTSIQSLGSDQSASAGSAGSVDHVSLSIPSATGPDVR 1732  
QY 58 ISEMFDLQEP--CLOTRLELY-KQGLRGLSLTKLKGPLTMASHYKHQC-PPTPETS--- 110  
DB 1733 TSSLSLSEQPTFTPMVVEAEIHASQGLSIPSTIGTETRVPPSPDSRPTSPSAGSH 1792  
QY 111 CATQIIITFESPKENLKDF 128  
DB 1793 TSVDLITFDSVAETLEDF 1810  
RESULT 9  
A56554  
transcription factor C/EBP - African clawed frog  
N;Alternate names: CCAAT/enhancer core binding protein  
C;Species: Xenopus laevis (African clawed frog)  
C;Date: 21-Jul-1995 #sequence\_revision 21-Jul-1995 #text\_change 09-Jul-2004  
C;Accession: A56554  
R;Xu, Q.; Tata, J.R.  
Mol. Cell. Dev. 38, 69-81, 1992  
A;Title: Characterization and developmental expression of Xenopus C/EBP gene.  
A;Reference number: A56554; MUID:9239265; PMID:1525039  
A;Accession: A56554  
A;Status: preliminary  
A;Molecule type: mRNA  
A;Residues: 1-305 <XUL>  
A;Cross-references: UNIPROT:Q91346; GB:S44193; NID:g255566; PIDN:AAE23276.1; PID:g255566  
A;Note: sequence extracted from NCBI backbone (NCBI:113707, NCBI:113708)  
C;Superfamily: CCAAT/enhancer-binding protein alpha  
C;Keywords: DNA binding; leucine zipper; transcription factor  
Query Match 10.6%; Score 81; DB 2; Length 305;  
Best Local Similarity 29.3%; Pred. No. 3.6;  
Matches 34; Conservative 17; Mismatches 35; Indels 30; Gaps 7;



A;Cross-references: UNIPROT.Q38897; GB:U39944; NID:g1122532; PIDN:AAB05099.1; PID:g1122532

C;Genetics:

A;Gene: BEL1

C;Superfamily: unassigned homeobox proteins; homeobox homology

C;Keywords: DNA binding; homeobox; nucleus; transcription regulation

F;331-450/domain: homeobox homology <HOX>

Query Match 10.3%; Score 79; DB 2; Length 610;  
Best Local Similarity 22.7%; Pred. No. 13;  
Matches 27; Conservative 20; Mismatches 42; Indels 30; Gaps 4;

QY 2 HHHHHSSGTEGRVAPARSPSTQPVEHVNAIQEARLLNLNRDTAAEMNE-----TV 55  
||||| : | : | : | : | : | : | : | :  
Db 72 HHHHHHQT-----SGGTQNQLLEDSSAMRLCNVNDFPSEVNDERPPQPS 119  
:  
QY 56 EVISEMEDLQEPCTCL-----QTRLELYKQGLRG-----SLTKLGPLTMMASHVKQH 102  
: | : | : | : | : | : | : | :  
Db 120 QGLSLSSLSSNNPTSLISQSFELRFQQOQQGGSGNKSTQHQLQHTQMNMNNSHHQNN 178  
:

RESULT 14

T45972

hypothetical protein F9D24.30 - Arabidopsis thaliana

C;Species: Arabidopsis thaliana (mouse-ear cress)

C;Date: 04-Feb-2000 #sequence\_revision 04-Feb-2000 #text\_change 09-Jul-2004

C;Accession: T45972

R;D'Angelo, M.; Vezzi, A.; Modesto, D.; Pigazzi, M.; Valle, G.; Mewes, H.W.; Lemcke, K.; submitted to the Protein Sequence Database, January 2000

A;Reference number: Z23011

A;Accession: T45972

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-329 <DAN>

A;Cross-references: UNIPROT.Q9M2K4; EMBL.AL137081

A;Experimental source: cultivar Columbia; BAC clone F9D24

C;Genetics:

A;Map position: 3

A;Introns: 208/2; 238/3; 281/1

A;Note: F9D24.30

Query Match 10.3%; Score 78.5; DB 2; Length 329;  
Best Local Similarity 28.0%; Pred. No. 6.8;  
Matches 21; Conservative 12; Mismatches 23; Indels 19; Gaps 3;

QY 2 HHHHHSSGTEGRVAPARSPSTQPVEHVNAIQE-----ARRLLNLSR 45  
||||| : | : | : | : | : | : | : | :  
Db 106 NHRRHS--INGNVPTRSSNTSTPSDH-NSLSDDDNNKEAPPDHDHMDNNVANQNN 162  
:

QY 46 DTAAMNETVEVISE 60  
||| : ||| :  
Db 163 AAGNNYNESDEVQSQ 177

RESULT 15

S25091

cruciferin BnC2 - rape

C;Species: Brassica napus (rape)

C;Date: 04-Feb-1998 #sequence\_revision 20-Feb-1998 #text\_change 09-Jul-2004

C;Accession: S25091

R;Green, J.P.; Crouch, M.L.  
Plant Mol. Biol. 19, 1049-1055, 1992

A;Title: Molecular analysis of a cruciferin storage protein gene family of Brassica napus

A;Reference number: S25090; MUID:92379259; PMID:1511129

A;Accession: S25091

A;Status: translation not shown

A;Molecule type: DNA

A;Residues: 1-496 <BRE>

A;Cross-references: UNIPROT.P33524; EMBL.X59295; NID:g17791; PIDN:CAA41985.1; PID:g76292

C;Genetics:

A;Gene: BnC2

A;Introns: 95/1; 222/2; 362/3

C;Superfamily: glycinin

C;Keywords: seed; storage protein

```

Query Match      10.2%; Score 78; DB 2; Length 496;
Best Local Similarity 23.1%; Pred. No. 13;
Matches 27; Conservative 19; Mismatches 51; Indels 20; Gaps 3;

Qy 19 RSPS-----PSTQPEHVNAIQEARLLNLSRDTAAEMNETVEISEMFDLQPTCL 70
   |::| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 219 RNFQFYLAGKNPQGSWLHGRQQPQNNILN-----GFPEVLAQAFKIDVRTAQ 269
   |::| | | | | | | | | | | | | | | | | | | | | | | | | | | |

Qy 71 QTRLELYKQGLRGLTKLKGPLTMASHYKQHCPTPTPETSATQIITFESFKENLKD 127
   |::| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 270 QLQ---NQQDNRGNIIVRVQPGFVIRPPLKSRPQETANGLEETICSARCTDNLDD 323
   |::| | | | | | | | | | | | | | | | | | | | | | | | | | | |

```

Search completed: March 8, 2005, 16:14:10  
Job time : 41 secs

GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: March 8, 2005, 15:56:25 ; Search time 177 Seconds  
(without alignments)  
410.821 Million cell updates/sec

Title: US-10-723-083-2  
Perfect score: 765  
Sequence: 1 MHHHHSHSGIEGRMAPARS.....ENLKDFLLVLPDCEWPEVQE 142

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 1612378 seqs, 512079187 residues

Total number of hits satisfying chosen parameters: 1612378

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Uniprot\_03.\*  
1: uniprot\_sprot.\*  
2: uniprot\_trembl.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	673	88.0	144	1 CSF2_HUMAN	P04141 homo sapien
2	664	86.8	144	2 Q647J8	Q647J8 homo sapien
3	653	85.4	144	2 Q9GL44	Q9GL44 macaca mula
4	633	82.7	144	2 Q865Y5	Q865Y5 papio anubi
5	548	71.6	144	1 CSF2_SHEEP	P28773 ovis aries
6	544	71.1	144	2 Q9MYK4	Q9MYK4 ovis aries
7	528	69.0	144	1 CSF2_CEREL	P51748 cervus elap
8	527	68.9	146	2 Q8WN17	Q8WN17 equus cabal
9	513	67.1	152	2 Q95L10	Q95L10 equus cabal
10	487	63.7	144	1 CSF2_PIG	Q29118 sus scrofa
11	486.5	63.6	143	2 Q6Q8A7	Q6Q8A7 bubalus bub
12	480.5	62.8	143	1 CSF2_BOVIN	P11052 bos taurus
13	473	61.8	144	1 CSF2_CANFA	P48749 canis famil
14	472	61.7	141	2 Q7YRF7	Q7YRF7 felis silve
15	451	59.0	144	1 CSF2_FELCA	Q62757 felis silve
16	441	57.6	127	1 CSF2_RAT	P48750 rattus norv
17	435.5	56.9	140	1 CSF2_CAVPO	Q60481 cavia porce
18	422.5	55.2	141	2 Q8VH40	Q8VH40 sigmodon hi
19	381.5	49.9	145	2 Q8CFB5	Q8CFB5 meriones un
20	361	47.2	138	2 Q99J91	Q99J91 marmota mon
21	360.5	47.1	141	1 CSF2_MOUSE	P01587 mus musculu
22	209.5	27.4	90	2 Q80XG1	Q80XG1 peromyscus
23	87	11.4	233	2 Q6ENY6	Q6ENY6 oryza sativ
24	87	11.4	499	2 Q81YK2	Q81YK2 homo sapien
25	85	11.1	354	2 Q8NDL5	Q8NDL5 homo sapien
26	85	11.1	499	2 Q8N7T5	Q8N7T5 homo sapien
27	84.5	11.0	196	1 HUNB_DROAA	Q46234 drosophila
28	84.5	11.0	2493	2 Q887B9	Q887B9 eastern equ
29	83	10.8	424	2 Q647M8	Q647M8 uncultured
30	83	10.8	785	2 Q8MQP8	Q8MQP8 drosophila
31	83	10.8	909	2 Q8THK7	Q8THK7 methanosaarc

32	82.5	10.8	610	2	Q9CS72	Q9CS72 mus musculu
33	81.5	10.7	344	2	Q8RYL3	Q8RYL3 oryza sativ
34	81.5	10.7	965	2	Q8JZS5	Q8JZS5 rattus norv
35	81.5	10.7	1212	2	Q8K4T4	Q8K4T4 rattus norv
36	81	10.6	171	1	HUNB_SCAAL	Q46254 scaptomyza
37	81	10.6	305	2	Q91346	Q91346 xenopus lae
38	81	10.6	343	2	Q9DG50	Q9DG50 xenopus lae
39	81	10.6	359	1	IKBB_MOUSE	Q60778 mus musculu
40	81	10.6	359	2	Q8VC27	Q8VC27 mus musculu
41	81	10.6	379	2	Q8SB73	Q8SB73 oryza sativ
42	81	10.6	379	2	Q7G6D7	Q7G6D7 oryza sativ
43	81	10.6	424	2	Q64E31	Q64E31 uncultured
44	80.5	10.5	285	2	Q8YN88	Q8YN88 anabaena sp
45	80.5	10.5	1371	2	Q7QRM3	Q7QRM3 giardia lam

ALIGNMENTS

RESULT 1  
ID CSF2\_HUMAN STANDARD; PRT; 144 AA.  
AC P04141: Q8NFI6;  
DT 01-NOV-1986 (Rel. 03, Created)  
DT 01-NOV-1986 (Rel. 03, Last sequence update)  
DE 25-OCT-2004 (Rel. 45, Last annotation update)  
DE Granulocyte-macrophage colony-stimulating factor precursor (GM-CSF)  
DE (Colony-stimulating factor) (CSF) (Sargramostim) (Molgramostin).  
GN Name=CSF2; Synonyms=GMCSF;  
OS Homo sapiens (Human)  
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
OX NCBI\_TaxID=9606;  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=85242684; PubMed=3925454;  
RA Lee F., Yokota T., Otsuka T., Gemmell L., Larson N., Luh J.,  
RA Arai K.-I., Rennick D.;  
RT "Isolation of cDNA for a human granulocyte-macrophage colony-  
stimulating factor by functional expression in mammalian cells.";  
RL Proc. Natl. Acad. Sci. U.S.A. 82:4360-4364(1985).  
RN [2]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=86205844; PubMed=3486413;  
RA Kaushansky K., O'Hara P.J., Berkner K., Segal G.M., Hagen F.S.,  
RA Adamson J.W.;  
RT "Genomic cloning, characterization, and multilineage growth-promoting  
activity of human granulocyte-macrophage colony-stimulating factor.";  
RL Proc. Natl. Acad. Sci. U.S.A. 83:3101-3105(1986).  
RN [3]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=85298329; PubMed=3898082;  
RA Cantrell M.A., Anderson D., Cerretti D.P., Price V., McKereghan K.,  
RA Tushinski R.J., Mochizuki D.Y., Larsen A., Grabstein K., Cosman D.;  
RT "Cloning, sequence, and expression of a human granulocyte/macrophage  
colony-stimulating factor.";  
RL Proc. Natl. Acad. Sci. U.S.A. 82:6250-6254(1985).  
RN [4]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=85218749; PubMed=3923623;  
RA Wong G.G., Wittek J.S., Temple P.A., Wilkens K.M., Leary A.C.,  
RA Luxenberg D.P., Jones S.S., Brown E.L., Kay R.M., Orr E.C.,  
RA Shoenaker C., Golde D.W., Kaufman R.J., Hewick R.M., Wang E.A.,  
RA Clark S.C.;  
RT "Human GM-CSF: molecular cloning of the complementary DNA and  
purification of the natural and recombinant proteins.";  
RL Science 228:810-815(1985).  
RN [5]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=86030234; PubMed=3876930;  
RA Miyake S., Otsuka T., Yokota T., Lee F., Arai K.-I.;  
RT "Structure of the chromosomal gene for granulocyte-macrophage colony  
stimulating factor: comparison of the mouse and human genes.";

EMBO J. 4:2561-2568(1985).  
 [6] SEQUENCE FROM N.A.  
 RP Kimmerly W., Bondoc M., Cheng J., Connolly K.S., Gunning K.M.,  
 RA Davis C.A., Kadner K., Migue T., Pitluck S., Pollard M., Rojeski H.,  
 RA Subramanian S., Martin C.H.;  
 RL Submitted (MAR-1998) to the EMBL/GenBank/DBJ databases.  
 [7] SEQUENCE FROM N.A., AND VARIANTS ILE-115 AND THR-117.  
 RP Riederer M.J., Carrington D.P., Chung M.-W., Lee C.L., Yi Q.,  
 RA Nickerson D.A.;  
 RL "SeattleSNPs: NHLBI HL66682 program for genomic applications, UW-  
 RT FHCRC, Seattle, WA (URL: <http://pga.gs.washington.edu>).";  
 RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.  
 [8] SEQUENCE OF 18-144 FROM N.A., AND VARIANT THR-117.  
 RP TISSUE=Peripheral blood;  
 RC Bhattacharya P., Pandey G., Mukherjee K.J.;  
 RA Submitted (MAY-2002) to the EMBL/GenBank/DBJ databases.  
 [9] CARBOHYDRATE-LINKAGE SITES.  
 RP MEDLINE=92144609; PubMed=1737041;  
 RX Kaushansky K., Lopez J.A., Brown C.B.;  
 RA "Role of carbohydrate modification in the production and secretion of  
 RT human granulocyte macrophage colony-stimulating factor in genetically  
 RT engineered and normal mesenchymal cells.";  
 RL Biochemistry 31:1881-1886(1992).  
 [10] X-RAY CRYSTALLOGRAPHY (2.4 ANGSTROMS).  
 RP MEDLINE=92108420; PubMed=1837174;  
 RX Dieterichs K., Boone T., Karplus P.A.;  
 RA "Novel fold and putative receptor binding site of granulocyte-  
 RT macrophage colony-stimulating factor.";  
 RL Science 254:1779-1782(1991).  
 [11] X-RAY CRYSTALLOGRAPHY (2.8 ANGSTROMS).  
 RP MEDLINE=92235844; PubMed=1569568;  
 RX Walter M.R., Cook W.J., Ballick S.E., Nagabhushan T.L., Trotta P.P.,  
 RA Bugg C.E.;  
 RT "Three-dimensional structure of recombinant human granulocyte-  
 RT macrophage colony-stimulating factor.";  
 RL J. Mol. Biol. 224:1075-1085(1992).  
 CC -I- FUNCTION: Cytokine that stimulates the growth and differentiation  
 CC of hematopoietic precursor cells from various lineages, including  
 CC granulocytes, macrophages, eosinophils and erythrocytes.  
 CC -I- SUBUNIT: Monomer.  
 CC -I- SUBCELLULAR LOCATION: Secreted.  
 CC -I- POLYMORPHISM: Variant Ile-117 may be a risk factor for atopic  
 CC asthma.  
 CC -I- PHARMACEUTICAL: Available under the names Leukine (Immunex) and  
 CC Leucomax (Novartis). Used in myeloid reconstitution following bone  
 CC marrow transplant, bone marrow transplant engraftment failure or  
 CC delay, mobilization and following transplantation of autologous  
 CC peripheral blood progenitor cells, and following induction  
 CC chemotherapy in older adults with acute myelogenous leukemia.  
 CC -I- SIMILARITY: Belongs to the GM-CSF family.  
 CC -I- DATABASE: NAME=Leukine; NOTE=Clinical information on Leukine;  
 CC WWW="http://www.leukine.com/".  
 -----  
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 DR EMBL; M13207; AAA98768.1; -;  
 DR EMBL; M11734; AAA52122.1; -;  
 DR EMBL; M11220; AAA52578.1; -;  
 DR EMBL; X03021; CAA26822.1; -;  
 DR EMBL; M10663; AAA52121.1; -;  
 DR EMBL; AAC04511; AAC08707.1; -;

DR EMBL; AF373868; AAKS1563.1; -;  
 DR EMBL; AF510855; AAM44054.1; -;  
 DR PIR; C24636; FOHUGM.  
 DR PDB; 1CSG; X-ray; A/B=18-144.  
 DR PDB; 2GMF; X-ray; A/B=-.  
 DR Genew; HGNC:2434; CSF2.  
 DR MIM; 138960; -;  
 DR GO; GO:0005129; F:granulocyte macrophage colony-stimulating f. . .; TAS.  
 DR InterPro; IPR009079; 4.helix\_cytokine.  
 DR InterPro; IPR000773; GM\_CSF.  
 DR Pfam; PF01109; GM\_CSF; 1.  
 DR PRINTS; PR00693; GMCSFACTOR.  
 DR ProDom; PD007349; GM\_CSF; 1.  
 DR SMART; SM00040; CSF2; 1.  
 DR PROSITE; PS00702; GM\_CSF; 1.  
 KW 3D-structure; Cytokine; Growth factor; Pharmaceutical;  
 KW Polymorphism; Signal.  
 FT SIGNAL 1 17  
 FT CHAIN 18 144  
 FT DISULFID 71 113  
 FT DISULFID 105 138  
 FT CARBOHYD 22 22  
 FT CARBOHYD 24 24  
 FT CARBOHYD 26 26  
 FT CARBOHYD 27 27  
 FT CARBOHYD 44 44  
 FT CARBOHYD 54 54  
 FT VARIANT 115 115  
 FT VARIANT 117 117  
 FT CONFLICT 96 96  
 FT CONFLICT 123 123  
 FT TURN 25 27  
 FT HELIX 30 45  
 FT HELIX 50 53  
 FT TURN 54 54  
 FT STRAND 56 60  
 FT TURN 66 67  
 FT HELIX 72 81  
 FT TURN 82 82  
 FT HELIX 85 90  
 FT HELIX 91 104  
 FT STRAND 115 119  
 FT HELIX 120 131  
 FT TURN 132 133  
 SQ SEQUENCE 144 AA; 16295 MW; 75D1E50506BCA7A8 CRC64;  
 Query Match 88.0%; Score 673; DB 1; Length 144;  
 Best Local Similarity 100.0%; Pred. No. 1.4e-54;  
 Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 16 APARSPSPSTQPEWHVNAIQEARRLLNLSRDAAENNETVEVISEMFDIQEPTCLQTRLE 75  
 Db 18 APARSPSPSTQPEWHVNAIQEARRLLNLSRDAAENNETVEVISEMFDIQEPTCLQTRLE 77  
 QY 76 LYKQGLRGSLLTKLKGPLTNWASHYKHCHCPPTETSCATQIITFESEPKENLKQFLVLPD 135  
 Db 78 LYKQGLRGSLLTKLKGPLTNWASHYKHCHCPPTETSCATQIITFESEPKENLKQFLVLPD 137  
 QY 136 CWEPVQE 142  
 Db 138 CWEPVQE 144  
 RESULT 2  
 Q647J8 PRELIMINARY; PRT; 144 AA.  
 ID Q647J8  
 AC Q647J8;  
 DT 25-OCT-2004 (TrEMBLrel. 28, Created)  
 DT 25-OCT-2004 (TrEMBLrel. 28, Last sequence update)  
 DT 25-OCT-2004 (TrEMBLrel. 28, Last annotation update)

[illegible]

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OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Caprinae; Ovis.
OX NCBI_TaxID=9940;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=92039044; PubMed=1937025; DOI=10.1016/0378-1119(91)90163-6;
RA McInnes C.J., Haig M.C.K.;
RT "Cloning and expression of a cDNA encoding ovine granulocyte-
macrophage colony-stimulating factor.";
RL Gene 105:275-279(1991).
CC -1- FUNCTION: Cytokine that stimulates the growth and differentiation
of hematopoietic precursor cells from various lineages, including
granulocytes, macrophages, eosinophils and erythrocytes (By
similarity).
CC -1- SUBUNIT: Monomer (By similarity).
CC -1- SUBCELLULAR LOCATION: Secreted.
CC -1- SIMILARITY: Belongs to the GM-CSF family.
CC -----
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CC -----
DR EMBL; X53561; CAA37632.1; -.
DR PIR; JH0469; JH0469.
DR HSSP; P04141; 2GMF.
DR InterPro; IPR009079; 4_helix_cytokine.
DR Pfam; PF01109; GM_CSF; 1.
DR PRINTS; PR00693; GMCSPFACTOR.
DR ProDom; PD007349; GM_CSF; 1.
DR SMART; SM00040; CSF2; 1.
DR PROSITE; PS00702; GM_CSF; 1.
DR SIGNAL 1 17 By similarity.
FT CHAIN 18 144 Granulocyte-macrophage colony-stimulating
factor.
FT DISULFID 71 113 By similarity.
FT DISULFID 105 138 By similarity.
FT CARBOHYD 44 44 N-linked (GlcNAc...) (Potential).
SQ SEQUENCE 144 AA; 16318 MW; ABAAC8733B580008 CRC64;

Query Match 71.6%; Score 548; DB 1; Length 144;
Best Local Similarity 80.3%; Pred. No. 5.7e-43;
Matches 102; Conservative 10; Mismatches 15; Indels 0; Gaps 0;

QY 16 APARSPSPQWEHVNIAQEARLLNLSRDTAAENNEIVEISEMFDLQEPICLQTRLE 75
Db 18 APTROPSPVTRPQHWDAIKEALSLNDSITDAVMDTEVVEVSEMFDSQEPICLQTRLE 77

QY 76 LYKQGLRGLSTLTKGLPLTMASHYKHCPPTPTSCATQIITFESPKENLKDFLLVIPP 135
Db 78 LYKQGLRGLSTLTKGLPLTMASHYKHCPPTPTSCATQIITFESPKENLKDFLLVIPP 137

QY 136 CWEPVQE 142
Db 138 CWEPVQK 144

RESULT 6
ID Q9MYK4 PRELIMINARY; PRT; 144 AA.
AC Q9MYK4;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Granulocyte-macrophage colony-stimulating factor (Fragment).
OS Ovis aries (Sheep).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;

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OC Caprinae; Ovis.
OX NCBI_TaxID=9940;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91331592; PubMed=1869289;
RA O'Brien P.M., Rothel J.S., Seow H.F., Wood P.R.;
RT "Cloning and sequencing of the cDNA for ovine granulocyte-Macrophage
colony stimulating factor (GM-CSF).";
RL Immunol. Cell Biol. 69:51-55(1991).
RN [2]
RP SEQUENCE FROM N.A.
RA O'Brien P.;
RL Submitted (OCT-1990) to the EMBL/GenBank/DBJ databases.
DR EMBL; X55991; CAA39463.1; -.
DR PIR; A61632; A61632.
DR HSSP; P04141; 2GMF.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005129; F:granulocyte macrophage colony-stimulating f...; IEA.
DR GO; GO:0006955; P:immune response; IEA.
DR InterPro; IPR009079; 4_helix_cytokine.
DR InterPro; IPR000773; GM_CSF.
DR Pfam; PF01109; GM_CSF; 1.
DR PRINTS; PR00693; GMCSPFACTOR.
DR ProDom; PD007349; GM_CSF; 1.
DR SMART; SM00040; CSF2; 1.
DR PROSITE; PS00702; GM_CSF; 1.
FT CHAIN 1 >144 granulocyte-macrophage colony-stimulating
factor.
FT NON_TER 144 144
SQ SEQUENCE 144 AA; 16290 MW; ABAAD7633B580008 CRC64;

Query Match 71.1%; Score 544; DB 2; Length 144;
Best Local Similarity 79.5%; Pred. No. 1.3e-42;
Matches 101; Conservative 10; Mismatches 16; Indels 0; Gaps 0;

QY 16 APARSPSPQWEHVNIAQEARLLNLSRDTAAENNEIVEISEMFDLQEPICLQTRLE 75
Db 18 APTROPSPVTRPQHWDAIKEALSLNDSITDAVMDTEVVEVSEMFDSQEPICLQTRLE 77

QY 76 LYKQGLRGLSTLTKGLPLTMASHYKHCPPTPTSCATQIITFESPKENLKDFLLVIPP 135
Db 78 LYKQGLRGLSTLTKGLPLTMASHYKHCPPTPTSCATQIITFESPKENLKDFLLVIPP 137

QY 136 CWEPVQE 142
Db 138 CWEPVQK 144

RESULT 7
ID CSF2 CEREL STANDARD; PRT; 144 AA.
AC P51748;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Granulocyte-macrophage colony-stimulating factor precursor (GM-CSF)
(Colony-stimulating factor) (CSF).
GN Name-CSF2;
OS Cervus elaphus (Red deer).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Cervidae;
OC Cervinae; Cervus.
OX NCBI_TaxID=9860;
RN [1]
RP SEQUENCE FROM N.A.
RA Lockhart E.A.;
RL Submitted (SEP-1994) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: Cytokine that stimulates the growth and differentiation
of hematopoietic precursor cells from various lineages, including
granulocytes, macrophages, eosinophils and erythrocytes (By
similarity).
CC -1- SUBUNIT: Monomer (By similarity).
CC -1- SUBCELLULAR LOCATION: Secreted.

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CC -I- SIMILARITY: Belongs to the GM-CSF family.
CC -----
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CC -----
DR EMBL; U14392; AAA21439.1; -.
DR HSSP; P04141; 2GMF.
DR InterPro; IPR009079; 4_helix_cytokine.
DR InterPro; IPR000773; GM-CSF.
DR Pfam; PF011109; GM-CSF; 1.
DR PRINTS; PR00693; GMCSFACTOR.
DR ProDom; PD007349; GM-CSF; 1.
DR SMART; SM00040; CSF2_1.
DR PROSITE; PS00702; GM-CSF; 1.
KW Cytokine; Glycoprotein; Growth factor; Signal.
FT SIGNAL 1 17
FT CHAIN 18 144
FT factor.
FT DISULFID 71 113
FT DISULFID 105 138
FT CARBOHYD 44 44
FT CARBOHYD 54 54
FT SEQUENCE 144 AA; 16283 MW; 1F5FFD03C94394 CRC64;
SQ
Query Match 59.0%; Score 528; DB 1; Length 144;
Best Local Similarity 77.2%; Pred. NO. 4.1e-41;
Matches 98; Conservative 10; Mismatches 19; Indels 0; Gaps 0;
QY 16 APARSPSPSTQPEWHVNAIQEARRLNLSRDTAAEMNETVEVISEMFDLQEPCLQTRLE 75
DB 18 APTRPSPVTRPQHVDAIKKLSLNSSDTRAAVNETVEVSENFDSQEPCLQTRLK 77
QY 76 LYKQGLRGLSLTKLKGPLTMASHYKQHCPTPTTSCATQIITPESKKNLKDFLLVIPP 135
DB 78 LYKQGLRGLSLTKLKGPLTMASHYKQHCPTPTTSCATQIITPESKKNLKDFLLVIPP 137
QY 136 CWEPVQE 142
DB 138 CWKPAQK 144
RESULT 8
Q8WN17 PRELIMINARY; PRT; 146 AA.
AC Q8WN17;
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Granulocyte-macrophage colony-stimulating factor (Fragment).
GN Name=GM-CSF;
OS Equus caballus (Horse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Perissodactyla; Equidae; Equus.
OX NCBI_TaxID=9796;
RN [1]
RP SEQUENCE FROM N.A.
RA Vecchione A., D'Mello F., Kanellos T.S., Howard C.R., Hamblin A.S.,
RL Submitted (AUG-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF448481; AAL41017.2; -.
DR HSSP; P04141; 2GMF.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005129; F:granulocyte macrophage colony-stimulating f. . .; IEA.
DR GO; GO:0006955; P:immune response; IEA.
DR Pfam; PF011109; GM-CSF; 1.
DR PRINTS; PR00693; GMCSFACTOR.
DR ProDom; PD007349; GM-CSF; 1.
DR SMART; SM00040; CSF2; 1.

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DR PROSITE; PS00702; GM-CSF; 1.
FT NON_TER 146 146
SQ SEQUENCE 146 AA; 16604 MW; B92D0617F391281C CRC64;
Query Match 68.9%; Score 527; DB 2; Length 146;
Best Local Similarity 77.2%; Pred. NO. 5.1e-41;
Matches 98; Conservative 13; Mismatches 16; Indels 0; Gaps 0;
QY 16 APARSPSPSTQPEWHVNAIQEARRLNLSRDTAAEMNETVEVISEMFDLQEPCLQTRLE 75
DB 18 APTRPSPVTRPQHVDAIKKLSLNSSDTRAAVNETVEVSENFDSQEPCLQTRLK 77
QY 76 LYKQGLRGLSLTKLKGPLTMASHYKQHCPTPTTSCATQIITPESKKNLKDFLLVIPP 135
DB 78 LYKQGLRGLSLTKLKGPLTMASHYKQHCPTPTTSCATQIITPESKKNLKDFLLVIPP 137
QY 136 CWEPVQE 142
DB 138 CWKPAQK 144
RESULT 9
Q95L10 PRELIMINARY; PRT; 152 AA.
AC Q95L10;
DT 01-DEC-2001 (TrEMBLrel. 19, Created)
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Granulocyte-macrophage colony-stimulating-factor.
GN Name=GM-CSF;
OS Equus caballus (Horse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Perissodactyla; Equidae; Equus.
OX NCBI_TaxID=9796;
RN [1]
RP SEQUENCE FROM N.A.
RA Maue S., Commandeur U., Steinbach F.;
RL Submitted (NOV-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY040203; AAK72108.2; -.
DR HSSP; P04141; 2GMF.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005129; F:granulocyte macrophage colony-stimulating f. . .; IEA.
DR GO; GO:0006955; P:immune response; IEA.
DR InterPro; IPR009079; 4_helix_cytokine.
DR InterPro; IPR000773; GM-CSF.
DR Pfam; PF011109; GM-CSF; 1.
DR PRINTS; PR00693; GMCSFACTOR.
DR ProDom; PD007349; GM-CSF; 1.
DR SMART; SM00040; CSF2; 1.
DR PROSITE; PS00702; GM-CSF; 1.
SQ SEQUENCE 152 AA; 17173 MW; 75605CC1ADE9EFE9 CRC64;
Query Match 67.1%; Score 513; DB 2; Length 152;
Best Local Similarity 78.7%; Pred. NO. 1.1e-39;
Matches 96; Conservative 11; Mismatches 15; Indels 0; Gaps 0;
QY 16 APARSPSPSTQPEWHVNAIQEARRLNLSRDTAAEMNETVEVISEMFDLQEPCLQTRLE 75
DB 18 APTRPSPVTRPQHVDAIKKLSLNSSDTRAAVNETVEVSENFDSQEPCLQTRLK 77
QY 76 LYKQGLRGLSLTKLKGPLTMASHYKQHCPTPTTSCATQIITPESKKNLKDFLLVIPP 135
DB 78 LYKQGLRGLSLTKLKGPLTMASHYKQHCPTPTTSCATQIITPESKKNLKDFLLVIPP 137
QY 136 CW 137
DB 138 CW 139
RESULT 10
CSF2_PIG STANDARD; PRT; 144 AA.
ID CSF2_PIG Q29046;
AC Q29118; Q29046;

```

DT 01-NOV-1997 (Rel. 35, Created)  
 DT 01-NOV-1997 (Rel. 35, Last sequence update)  
 DT 05-JUL-2004 (Rel. 44, Last annotation update)  
 DE Granulocyte-macrophage colony-stimulating factor precursor (GM-CSF)  
 DE (Colony-stimulating factor) (CSF).  
 GN Name=CSF2;  
 OS Sus scrofa (Pig).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.  
 OX NCBI\_TaxID=9823;  
 RN [1]  
 RN SEQUENCE FROM N.A.  
 RC TISSUE=Spleen;  
 RA Foss D.L., Murtaugh M.P.;  
 RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.  
 RN [2]  
 RN SEQUENCE FROM N.A.  
 RA Gloster S.E., Sandeman R.M., Strom A.D.G.;  
 RL Submitted (AUG-1996) to the EMBL/GenBank/DBJ databases.  
 RN [3]  
 RN SEQUENCE FROM N.A.  
 RX MEDLINE=96167041; PubMed=8595928;  
 RA Inumaru S., Takamatsu H.;  
 RT "CDNA cloning of porcine granulocyte-macrophage colony-stimulating  
 factor.";  
 RL Immunol. Cell Biol. 73:474-476(1995).  
 RN [4]  
 RN SEQUENCE FROM N.A.  
 RA Cho Y.W., Choi I.-S., Yoo H.S.;  
 RT "Cloning of porcine granulocyte macrophage-colony stimulating factor  
 in alveolar macrophages.";  
 RL Submitted (MAY-2002) to the EMBL/GenBank/DBJ databases.  
 CC -I- FUNCTION: Cytokine that stimulates the growth and differentiation  
 CC of hematopoietic precursor cells from various lineages, including  
 CC granulocytes, macrophages, eosinophils and erythrocytes (by  
 CC similarity).  
 CC -I- SUBUNIT: Monomer.  
 CC -I- SUBCELLULAR LOCATION: Secreted.  
 CC -I- SIMILARITY: Belongs to the GM-CSF family.  
 CC  
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 CC or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch)).  
 CC  
 DR EMBL; U61139; AAB03867.1; -;  
 DR EMBL; U67318; AAB49939.1; -;  
 DR EMBL; U67175; AAB06854.1; -;  
 DR EMBL; D21074; BAA04649.1; -;  
 DR EMBL; AV116504; AAM48280.1; -;  
 DR HSSP; P04141; 2GMF.  
 DR InterPro; IPR009079; 4\_helix\_cytokine.  
 DR InterPro; IPR000773; GM-CSF.  
 DR Pfam; PF01109; GM-CSF; 1.  
 DR PRINTS; PR00693; GMCSFACTOR.  
 DR ProDom; PD007349; GM-CSF; 1.  
 DR SMART; SM00040; CSF2; 1.  
 DR PROSITE; PS00702; GM-CSF; 1.  
 DR Cytokine; Glycoprotein; Growth factor; Signal.  
 FT SIGNAL 1 17 Potential.  
 FT CHAIN 18 144 Granulocyte-macrophage colony-stimulating  
 FT factor.  
 FT FT 71 113 By similarity.  
 FT DISULFID 105 138 By similarity.  
 FT CARBOHYD 44 44 N-linked (GlcNAc...) (Potential).  
 FT CARBOHYD 47 47 N-linked (GlcNAc...) (Potential).  
 FT CARBOHYD 54 54 N-linked (GlcNAc...) (Potential).  
 FT CONFLICT 59 59 V -> I (in Ref. 2).  
 FT CONFLICT 140 140 G -> E (in Ref. 2).  
 FT CONFLICT 142 143 VK -> AQ (in Ref. 2).

SQ SEQUENCE 144 AA; 16254 MW; 793DACB62CF736D0 CRC64;  
 Query Match 63.7%; Score 487; DB 1; Length 144;  
 Best Local Similarity 70.1%; Pred. No. 2.6e-37;  
 Matches 89; Conservative 16; Mismatches 22; Indels 0; Gaps 0;  
 QY 16 APARSPSPSTQPEWHVNAIQEARRLLNLSRDFAAEMNETVEVISEMFDLQEPCTCLQTRLE 75  
 DB 18 ATRPPSPVTRPQVHQVDAIKEALSLNNSNDFAAVNNEIVDVVCEMFDQEPCTCLQTRLN 77  
 QY 76 LYKQGLRGLSLTKLKGPLTMASHYKQHCPTTSCATQIITFESFKENLKDPLLVIPFD 135  
 DB 78 LYKQGLRGLSLTKLKSPLTLAKHYEQHCPLTTESTCETQSIITFKSFKDSLKNKFLFIIPFD 137  
 QY 136 CWEPVQVE 142  
 DB 138 CWGPVKK 144  
 RESULT 11  
 Q6Q8A7 PRELIMINARY; PRT; 143 AA.  
 AC Q6Q8A7;  
 DT 05-JUL-2004 (TReMBLrel. 27, Created)  
 DT 05-JUL-2004 (TReMBLrel. 27, Last sequence update)  
 DT 05-JUL-2004 (TReMBLrel. 27, Last annotation update)  
 DE Granulocyte macrophage colony stimulating factor.  
 OS Bubalus bubalis (Domestic water buffalo).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
 OC Bovinae; Bubalus.  
 OX NCBI\_TaxID=89462;  
 RN [1]  
 RN SEQUENCE FROM N.A.  
 RA Dhinakar Raj G., HariShankar M., Thennarasu S., Mahalinga Nainar A.;  
 RL Submitted (FEB-2004) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AY553190; AAS59070.1; -;  
 DR GO; GO:0005129; R:granulocyte macrophage colony-stimulating f...; IEA.  
 DR GO; GO:0005955; P:immune response; IEA.  
 DR InterPro; IPR009079; 4\_helix\_cytokine.  
 DR InterPro; IPR000773; GM-CSF.  
 DR Pfam; PF01109; GM-CSF; 1.  
 DR PRINTS; PR00693; GMCSFACTOR.  
 DR ProDom; PD007349; GM-CSF; 1.  
 DR SMART; SM00040; CSF2; 1.  
 DR PROSITE; PS00702; GM-CSF; 1.  
 SQ SEQUENCE 143 AA; 16105 MW; 274532FF0F0FE4C3F CRC64;  
 Query Match 63.6%; Score 486.5; DB 2; Length 143;  
 Best Local Similarity 70.1%; Pred. No. 2.9e-37;  
 Matches 89; Conservative 18; Mismatches 19; Indels 1; Gaps 1;  
 QY 16 APARSPSPSTQPEWHVNAIQEARRLLNLSRDFAAEMNETVEVISEMFDLQEPCTCLQTRLE 75  
 DB 18 ATRPPSPVTRPQVHQVDAIKEALSLNNSNDFAAVNNEIVDVVCEMFDQEPCTCLQTRLK 76  
 QY 76 LYKQGLRGLSLTKLKGPLTMASHYKQHCPTTSCATQIITFESFKENLKDPLLVIPFD 135  
 DB 77 LYKQGLRGLSLTKLMSGLTMATHYKHCPTTSCGTQITFKSFKEDLKEFLFIIPFD 136  
 QY 136 CWEPVQVE 142  
 DB 137 CWEPQAK 143  
 RESULT 12  
 CSF2\_BOVIN STANDARD; PRT; 143 AA.  
 ID CSF2\_BOVIN  
 AC P11052;  
 DT 01-JUL-1989 (Rel. 11, Created)  
 DT 01-JUL-1989 (Rel. 11, Last sequence update)  
 DT 05-JUL-2004 (Rel. 44, Last annotation update)

DE Granulocyte-macrophage colony-stimulating factor precursor (GM-CSF)  
 DE (Colony-stimulating factor) (CSF).  
 GN Name=CSF2;  
 OS Bos taurus (Bovine).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
 OC Bovinae; Bos.  
 OX NCBI\_TaxID=9913;  
 RN [1]  
 RN SEQUENCE FROM N.A.  
 RP MEDLINE=89096971; PubMed=3062386; DOI=10.1016/0161-5890(88)90120-4;  
 RX Mallatzevski C.R., Schoenborn M.A., Cerretti D.P., Wignall J.M.,  
 RA Picha K.S., Cosman D., Tushinski R.J., Gillis S., Baker P.E.,  
 RT "Bovine GM-CSF: molecular cloning and biological activity of the  
 RL recombinant protein.";  
 RL Mol. Immunol. 25:843-850(1988).  
 RN [2]  
 RN SEQUENCE FROM N.A.  
 RP MEDLINE=90021093; PubMed=2678728; DOI=10.1016/0165-2427(89)90036-6;  
 RX Leong S.K., Flagg G.M., Lawman M.J.P., Gray P.W.;  
 RA "Cloning and expression of the cDNA for bovine granulocyte-macrophage  
 RT colony-stimulating factor.";  
 RL Vet. Immunol. Immunopathol. 21:261-278(1989).  
 CC -!- FUNCTION: Cytokine that stimulates the growth and differentiation  
 CC of hematopoietic precursor cells from various lineages, including  
 CC granulocytes, macrophages, eosinophils and erythrocytes (By  
 CC similarity).  
 CC -!- SUBUNIT: Monomer (By similarity).  
 CC -!- SUBCELLULAR LOCATION: Secreted.  
 CC -!- SIMILARITY: Belongs to the GM-CSF family.  
 CC  
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 CC  
 DR EMBL; U22385; AAA66075.1; -;  
 DR PIR; J00037; F0BOGM.  
 DR HSSP; P04141; 2GMF.  
 DR InterPro; IPR009079; 4 helix\_cytokine.  
 DR InterPro; IPR000773; GM\_CSF.  
 DR Pfam; PF01109; GM\_CSF; 1.  
 DR PRINTS; PR00693; GMCSPACTOR.  
 DR ProDom; PD007349; GM\_CSF; 1.  
 DR SMART; SM00040; CSF2; 1.  
 DR PROSITE; PS00702; GM\_CSF; 1.  
 DR Cytokine; Glycoprotein; Growth factor; Signal.  
 FT SIGNAL 1 17  
 FT CHAIN 18 143 Granulocyte-macrophage colony-stimulating  
 FT factor.  
 FT By similarity.  
 FT DISULFID 70 112 By similarity.  
 FT DISULFID 104 137 By similarity.  
 FT CARBOHYD 44 44 N-linked (GlcNAc...) (Potential).  
 FT CARBOHYD 54 54 N-linked (GlcNAc...) (Potential).  
 SQ SEQUENCE 143 AA; 16157 MW; 4A24E26A870A51BC CRC64;  
 Query Match 62.8%; Score 480.5; DB 1; Length 143;  
 Best Local Similarity 68.5%; Pred. No. 1e-36;  
 Matches 87; Conservative 20; Mismatches 19; Indels 1; Gaps 1;  
 QY 16 APARSPSPQPEHVNVAIQEARRLLNLSRDTAENNETVEVISEMFDLOEPTCLOTRLR 75  
 DB 18 APTRPNTATRPQWDAIQEALLSLNHSDDTAVMMDT-EVSEKFDSEPTCLOTRLK 76  
 QY 76 LYKQGLRGSITLKGPLTMASHYKQHCPTPTSCATQIITPESFKENLKDFLLVPPD 135  
 DB 77 LYKNGLSITSLMGLSITMTATHYKHCPTPTSCGTQISFNKFNKDFLLFIIPD 136  
 QY 136 CWEPVQE 142  
 |||||

Db 137 CWEPQAK 143  
 RESULT 13  
 CSF2 CANFA  
 ID \_CSF2 CANFA STANDARD; PRT; 144 AA.  
 AC P48749;  
 DT 01-FEB-1996 (Rel. 33, Created)  
 DT 01-FEB-1996 (Rel. 33, Last sequence update)  
 DT 05-JUL-2004 (Rel. 44, Last annotation update)  
 DE Granulocyte-macrophage colony-stimulating factor precursor (GM-CSF)  
 DE (Colony-stimulating factor) (CSF).  
 GN Name=CSF2;  
 OS Canis familiaris (Dog).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.  
 OX NCBI\_TaxID=9615;  
 RN [1]  
 RN SEQUENCE FROM N.A.  
 RP MEDLINE=91329842; PubMed=1868252;  
 RX Nash R.A., Schueniger F., Appelbaum F.R., Hammond W.P., Boone T.,  
 RA Morris C.F., Slichter S.J., Storb R.;  
 RT "Molecular cloning and in vivo evaluation of canine granulocyte-  
 RL macrophage colony-stimulating factor.";  
 RL Blood 78:930-937(1991).  
 CC -!- FUNCTION: Cytokine that stimulates the growth and differentiation  
 CC of hematopoietic precursor cells from various lineages, including  
 CC granulocytes, macrophages, eosinophils and erythrocytes (By  
 CC similarity).  
 CC -!- SUBUNIT: Monomer (By similarity).  
 CC -!- SUBCELLULAR LOCATION: Secreted.  
 CC -!- SIMILARITY: Belongs to the GM-CSF family.  
 CC  
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 CC  
 DR EMBL; S49738; AAB19466.1; -;  
 DR PIR; A4936; A4936.  
 DR HSSP; P04141; 2GMF.  
 DR InterPro; IPR009079; 4 helix\_cytokine.  
 DR InterPro; IPR000773; GM\_CSF.  
 DR Pfam; PF01109; GM\_CSF; 1.  
 DR PRINTS; PR00693; GMCSPACTOR.  
 DR ProDom; PD007349; GM\_CSF; 1.  
 DR SMART; SM00040; CSF2; 1.  
 DR PROSITE; PS00702; GM\_CSF; 1.  
 DR Cytokine; Glycoprotein; Growth factor; Signal.  
 FT SIGNAL 1 17 By similarity.  
 FT CHAIN 18 144 Granulocyte-macrophage colony-stimulating  
 FT factor.  
 FT By similarity.  
 FT DISULFID 71 113 By similarity.  
 FT DISULFID 105 138 By similarity.  
 FT CARBOHYD 44 44 N-linked (GlcNAc...) (Potential).  
 SQ SEQUENCE 144 AA; 16137 MW; 6323807E1F6C343 CRC64;  
 Query Match 61.8%; Score 473; DB 1; Length 144;  
 Best Local Similarity 68.5%; Pred. No. 5.2e-36;  
 Matches 87; Conservative 19; Mismatches 21; Indels 0; Gaps 0;  
 QY 16 APARSPSPQPEHVNVAIQEARRLLNLSRDTAENNETVEVISEMFDLOEPTCLOTRLR 75  
 DB 18 APTRPNTATRPQWDAIQEALLSLNHSDDTAVMMDT-EVSEKFDSEPTCLOTRLK 77  
 QY 76 LYKQGLRGSITLKGPLTMASHYKQHCPTPTSCATQIITPESFKENLKDFLLVPPD 135  
 DB 78 LYKEGLQGSITSLKNPLTMMANHYKQHCPTPTSPCATQINFKSPKENLKDFLENIIPD 137  
 QY 136 CWEPVQE 142  
 |||||



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OM protein - protein search, using sw model

Run on: March 8, 2005, 15:55:38 ; Search time 164 Seconds  
(without alignments)  
334.878 Million cell updates/sec

Title: us-10-723-083-2

Perfect score: 765

Sequence: 1 MHHHHSSGIEGRMAPARS.....ENLKDFLLVLPDCWEPVOE 142

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues

Total number of hits satisfying chosen parameters: 2105692

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A\_Geneseq\_16Dec04:\*

- 1: Geneseqp1980s:\*
- 2: Geneseqp1990s:\*
- 3: Geneseqp2000s:\*
- 4: Geneseqp2001s:\*
- 5: Geneseqp2002s:\*
- 6: Geneseqp2003as:\*
- 7: Geneseqp2003bs:\*
- 8: Geneseqp2004s:\*

Pred' No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	682	89.2	259	2	AAR79317 IL-3 cont
2	682	89.2	259	3	AAY53198 Human int
3	682	89.2	259	4	AAEI3992 Myelopoie
4	682	89.2	259	5	ABG97765 Human int
5	682	89.2	259	8	ADJ14353 Chimera p
6	679	88.8	533	2	AAW19763 p53-GW-CS
7	678	88.6	128	1	AAP90118 Synthetic
8	678	88.6	128	1	AAP90115 Synthetic
9	678	88.6	128	2	AAR79338 pMON13012
10	678	88.6	128	2	AAW00103 Granulocy
11	678	88.6	128	3	AAAY53217 Human G-C
12	678	88.6	128	4	AAEI4011 Chemical
13	678	88.6	128	5	ABG97784 Human int
14	678	88.6	128	8	ADJ14372 Protein r
15	678	88.6	274	2	AAR79320 IL-3 cont
16	678	88.6	274	3	AAAY53201 Human int
17	678	88.6	274	4	AAEI3995 Myelopoie
18	678	88.6	274	5	ABG97768 Human int
19	678	88.6	274	8	ADJ14356 Chimera p
20	678	88.6	301	2	AAR79318 IL-3 cont
21	678	88.6	301	3	AAAY53199 Human int
22	678	88.6	301	4	AAEI3993 Chemical
23	678	88.6	301	5	ABG97766 Human int
24	676	88.4	523	3	AAAY44994 HD70scFv-
25	675	88.2	712	8	ADL16720 Human stu

## ALIGNMENTS

## RESULT 1

AAAR79317  
ID AAR79317 standard; protein; 259 AA.

XX AAR79317;

XX AC

XX XX

DT 25-AUG-1999 (first entry)

XX XX

DE IL-3 containing fusion protein.

XX XX

KW interleukin; hIL-3; CSF; colony stimulating factor; cytokine; lymphokine;

KW mutant; mutein; fusion protein.

XX Synthetic.

XX WO9521254-A1.

XX PD 10-AUG-1995.

XX PF 02-FEB-1995; 95WO-US001185.

XX PR 04-FEB-1994; 94US-00192325.

XX (SEAR ) SEARLE & CO G D.

XX Bauer CS, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;

XX Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;

XX WPI; 1995-283774/37.

XX N-PSDB; AAQ97169.

XX Fusion proteins comprising a human interleukin-3 variant, a linker and

XX interleukin-3, a variant or a colony stimulating factor - useful to

XX increase haematopoietic cell prodn. in a mammal.

XX Claim 16; Page 86; 447pp; English.

XX A new fusion protein has the formula R1-L-R2, R2-L-R1, R1-R2, R2-R1, R1-L

XX (hIL-3) having the present generic sequence, R2 is a second colony

XX stimulating factor (CSF) including cytokine, lymphokine, interleukin,

XX haematopoietic growth factor or IL-3 variant, and L is a linker. The

XX present sequence corresponds to native hIL-3(1-133) in which 1-14 amino

XX acids may be deleted from the N-terminal, 1-15 amino acids can be deleted

XX from the C-terminal, and at least 4 and up to 44 amino acids in the

XX region 17-123 are different from those in the native protein. The fusion

XX protein is used to increase haematopoietic cell production. It is also

XX useful as an IL-3 antagonist or as a discrete antigenic fragment for

XX

XX

XX

XX

XX

XX

XX

XX

XX

CC production of antibodies useful in immunoassays and immunotherapy.  
 CC Antagonists are used to block the growth of certain cancer cells and in  
 CC treatment of asthma. The fusion protein can also be used to stimulate  
 CC bone marrow and blood cell activation and growth in vitro before infusion  
 CC , and to treat diseases characterised by decreased levels of myeloid,  
 CC erythroid, lymphoid and/or megakaryocyte cells of the haematopoietic  
 CC system. The protein has the usual activity of both its component  
 CC proteins, but may have increased synergistic activity and reduced  
 CC undesired side effects  
 XX  
 SQ Sequence 259 AA;

Query Match 89.2%; Score 682; DB 2; Length 259;  
 Best Local Similarity 97.0%; Pred. No. 2.2e-64;  
 Matches 130; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 9 SGIEGRMAPRSPSTQPEWHNAIQEARRLLNLSRDAAENNEVEISEMFDLQEPT 68  
 DB 126 SGGGSNMAPRSPSTQPEWHNAIQEARRLLNLSRDAAENNEVEISEMFDLQEPT 185  
 QY 69 CLQTRLELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLQDF 128  
 DB 186 CLQTRLELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLQDF 245  
 QY 129 LLVIPDCWEPVQE 142  
 DB 246 LLVIPDCWEPVQE 259

RESULT 2  
 AA53198  
 ID AAY53198 standard; protein; 259 AA.  
 AC AAY53198;  
 DT 19-MAY-2000 (first entry)  
 XX Human interleukin-3 mutant containing fusion protein SEQ ID NO:141.  
 DE Human; interleukin 3; IL-3; mutant; mutein; CSF; cytokine;  
 KW colony stimulating factor; haematopoietic growth factor; lymphokine;  
 KW fusion protein; haematopoietic disorder; infection; cancer;  
 KW radiation therapy; chemotherapy; bone marrow suppressive drug;  
 KW bone marrow activation; blood cell activation; blood transplant.  
 XX Homo sapiens.  
 OS Synthetic.  
 OS  
 XX US6022535-A.  
 XX 08-FEB-2000.  
 XX 06-JUN-1995; 95US-00469318.  
 XX 04-FEB-1994; 94US-00192325.  
 PR 02-FEB-1995; 95WO-US001185.  
 PR 06-APR-1995; 95US-00411795.  
 XX (SEAR ) SEARLE & CO G D.  
 XX Bauer SC, Abrams MA, Braford-Goldberg SR, Easton AM, Klein BK;  
 PI Paik K, Thomas JW, McKearn JP, Olins PO, Caparon MH;  
 XX WPI; 2000-160368/14.  
 XX Treating hematopoietic disorders with fusion proteins comprising mutated  
 PT interleukin-3 fused with secondary colony stimulating factors or other  
 PT interleukin-3 variants.  
 XX Example 27; Col 71-72; 276pp; English.  
 PS Methods have been developed for treating haematopoietic disorders with  
 XX fusion proteins comprising recombinant, mutated human interleukin-3 (hIL-

CC 3) variants or mutant proteins (muteins) fused with secondary colony  
 CC stimulating factors (CSFs) (e.g. cytokines, lymphokines, interleukin  
 CC and/or haematopoietic colony stimulating factors) or other interleukin-3  
 CC variants with or without a linker. The methods may be used in vivo to  
 CC treat haematopoietic disorders resulting from bacterial, viral and fungal  
 CC infections, cancer radiation therapy, chemotherapy or bone marrow  
 CC suppressive drugs. They may also be used in vitro to stimulate bone  
 CC marrow and blood cell activation and growth prior to infusion of the bone  
 CC marrow and blood transplants into patients. IL-3 is a haematopoietic  
 CC growth factor which has the property of being able to promote the  
 CC survival, growth and differentiation of haematopoietic cells. The fusion  
 CC molecules are characterised by possessing the usual activity of both of  
 CC their constituent peptides and further by having a biological or  
 CC physiological activity greater than the additive function of the IL-3 or  
 CC second CSF alone (i.e. the peptides act synergistically). Their activity  
 CC may also be further enhanced by the mutations they comprise. The  
 CC variations may further reduce undesirable side effects associated with IL  
 CC -3. AAY53130 to AAY53226, and AAA03721 to AAA03782 represent sequences  
 CC used in the exemplification of the present invention  
 XX  
 SQ Sequence 259 AA;

Query Match 89.2%; Score 682; DB 3; Length 259;  
 Best Local Similarity 97.0%; Pred. No. 2.2e-64;  
 Matches 130; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 9 SGIEGRMAPRSPSTQPEWHNAIQEARRLLNLSRDAAENNEVEISEMFDLQEPT 68  
 DB 126 SGGGSNMAPRSPSTQPEWHNAIQEARRLLNLSRDAAENNEVEISEMFDLQEPT 185  
 QY 69 CLQTRLELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLQDF 128  
 DB 186 CLQTRLELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLQDF 245  
 QY 129 LLVIPDCWEPVQE 142  
 DB 246 LLVIPDCWEPVQE 259

RESULT 3  
 AA531992  
 ID AAE13992 standard; protein; 259 AA.  
 XX AAE13992;  
 XX 26-FEB-2002 (first entry)  
 DT Myelopietin (MPO) protein #127.  
 XX Myelopietin conjugate; MPO; immunosuppressive; vulnary; antiparasitic;  
 KW antibacterial; virucide; interleukin-3; IL-3; haematopoietic disorder;  
 KW haematopoietic growth factor receptor; neutropenia; thrombocytopenia;  
 KW leukopenia; anaemia; chemotherapy; bone marrow transplantation; burn;  
 KW radiation therapy; haematopoietic progenitor mobilisation; infection;  
 XX wound healing.  
 XX Unidentified.  
 XX WO200176639-A2.  
 XX 18-OCT-2001.  
 XX 06-APR-2001; 2001WO-US011256.  
 XX 06-APR-2000; 2000US-0195496P.  
 XX (PHAA ) PHARMACIA CORP.  
 PA (FINN/) FINN R.  
 PA (GOKA/) GOKARN Y.  
 PA (HILL/) HILLS R.  
 PA (NICA/) NICASTRO P.  
 PA (QIHH/) QI H.  
 PA (SEDO/) SEDO K.



PA (SIEGEL) SIEGEL N.  
 PA (WALT) WALTER S.  
 XX Finn R, Gokarn Y, Hills R, Nicastro P, Qi H, Sedo K, Siegel N;  
 PI Walter S;  
 XX WPI; 2001-657130/75.  
 XX Myelopoiectin conjugate for treating e.g. leukopenia comprises a water-  
 PT soluble polymer attached to the protein.  
 XX Claim 5; Page 197-198; 429pp; English.  
 XX The invention relates to chemically modified myelopoiectin (MPO)  
 CC conjugates having at least one water-soluble polymer molecule covalently  
 CC attached, via activating group, to at least one amino acid residue of a  
 CC biologically active myelopoiectin polypeptide. MPOs are multifunctional  
 CC agonists of human interleukin-3 (IL-3) and another haematopoietic growth  
 CC factor receptor. Sequences of the invention are useful for treating  
 CC haematopoietic disorders (e.g. neutropenia, leukopenia, thrombocytopenia  
 CC and anaemia), including those arising from chemotherapy and radiation  
 CC therapy. MPOs are also useful in bone marrow transplantation, wound  
 CC healing, burn treatment and the treatment of parasite, bacterial or viral  
 CC infection. They are useful for mobilising haematopoietic progenitors and  
 CC stem cells. The chemically modified MPOs have a longer lasting neutrophil  
 CC -releasing activity, decreased clearance rate, increased stability and  
 CC decreased antigenicity than unmodified myelopoiectins. The present  
 CC sequence is a myelopoiectin protein  
 XX Sequence 259 AA;  
 SQ  
 Query Match 89.2%; Score 682; DB 4; Length 259;  
 Best Local Similarity 97.0%; Pred. No. 2.2e-64;  
 Matches 130; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 9 SGIEGMAPARSPSPSTQPMWEHVNATQEARLLNLSRDTAENNETVEVISEMFDLOEPT 68  
 DB 126 SGGGNNAPARSPSPSTQPMWEHVNATQEARLLNLSRDTAENNETVEVISEMFDLOEPT 185  
 QY 69 CLOTRLELYKQGLRGSITLKLKGLPTWASHYKHQCPTPTTSCATQIITPESKFNKDF 128  
 DB 186 CLOTRLELYKQGLRGSITLKLKGLPTWASHYKHQCPTPTTSCATQIITPESKFNKDF 245  
 QY 129 LLVIPDCWEPVQES 142  
 DB 246 LLVIPDCWEPVQES 259  
 RESULT 4  
 ABG97765  
 ID ABG97765 standard; protein; 259 AA.  
 XX  
 AC ABG97765;  
 XX  
 DT 18-DEC-2002 (first entry)  
 XX  
 DE Human Interleukin-3 chimaeric protein #20.  
 DE  
 DE Human; interleukin-3; IL-3; mutant; mutagen; stem cell;  
 KW haematopoietic factor; GM-CSF; colony stimulating factor; CSF-1; G-CSF;  
 KW G-CSFser17; c-mpl ligand; TPO; MGDF; erythropoietin; flt3 ligand;  
 KW human growth hormone; B-cell growth factor; leukaemia;  
 KW B-cell differentiation factor; eosinophil differentiation factor;  
 KW stem cell factor; SCF; cyclic neutropenia; aplastic anaemia;  
 KW thrombocytopenia; idiopathic neutropenia; Chediak-Higashi syndrome;  
 KW systemic lupus erythematosus; SLE; myelodysplastic syndrome;  
 KW myelofibrosis.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 OS  
 XX  
 PN US6436387-B1.  
 XX

PD 20-AUG-2002.  
 XX  
 PF 09-DEC-1996; 96US-00762227.  
 XX  
 PR 24-NOV-1992; 92US-00981044.  
 XX  
 PR 22-NOV-1993; 93WO-US011197.  
 XX  
 PR 04-FEB-1994; 94US-00192325.  
 XX  
 PR 04-FEB-1995; 95WO-US001185.  
 XX  
 PR 06-APR-1995; 95US-00411795.  
 XX  
 PR 06-JUN-1995; 95US-00446872.  
 XX  
 PA (SEAR) SEARLE & CO G D.  
 XX  
 XX Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
 PI Klein BK, McKearn JP, Ollins PO, Paik K, Thomas JW;  
 XX WPI; 2002-749206/81.  
 XX  
 XX Ex vivo expansion of stem cells, for enhancing transduction efficiency of  
 PT cultured stem cells, comprises culturing stem cells in growth medium  
 PT having mutant interleukin-3, and hematopoietic factor, and harvesting  
 PT cultured cells.  
 XX  
 PS Claim 8; Col 247-250; 203pp; English.  
 XX  
 CC The invention relates to ex vivo expansion of stem cells, comprises  
 CC culturing stem cells with a growth medium comprising a chimaera protein,  
 CC and harvesting the cultured stem cells. The chimaera is based on a  
 CC mutated human interleukin-3 (IL-3) sequence coupled to a haematopoietic  
 CC factor (e.g. GM-CSF (colony stimulating factor), CSF-1, G-CSF, G-  
 CC CSFser17, c-mpl ligand TPO, MGDF, erythropoietin, IL-1-13, IL-15, IL-16,  
 CC flt3 ligand, human growth hormone, B-cell growth factor, B-cell  
 CC differentiation factor, eosinophil differentiation factor and stem cell  
 CC factor (SCF)) via a peptide linker. The formula for the chimaera is given  
 CC in the specification. Also included is a method for enhancing the  
 CC efficiency of the transduction of cultured stem cells by a heterologous  
 CC gene, comprising: (a) removing stem cells from a patient or donor; (b)  
 CC culturing the stem cells with a growth medium comprising the chimaera (c)  
 CC transducing DNA into cultured cells; and (d) harvesting the transduced  
 CC cells. The method is useful for ex vivo expansion of stem cells, and  
 CC enhancing the efficiency of the transduction of cultured stem cells by a  
 CC heterologous gene. The method is also useful for treating a patient  
 CC having a haematopoietic disorder. The expanded haematopoietic cells are  
 CC also useful in the treatment of cyclic neutropenia, aplastic anaemia,  
 CC thrombocytopenia, idiopathic neutropenia, Chediak-Higashi syndrome,  
 CC systemic lupus erythematosus (SLE), leukaemia, myelodysplastic syndrome  
 CC and myelofibrosis. The present sequence is a human IL-3  
 CC mutant/haematopoietic factor chimaeric sequence  
 XX  
 SQ Sequence 259 AA;  
 Query Match 89.2%; Score 682; DB 5; Length 259;  
 Best Local Similarity 97.0%; Pred. No. 2.2e-64;  
 Matches 130; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
 QY 9 SGIEGMAPARSPSPSTQPMWEHVNATQEARLLNLSRDTAENNETVEVISEMFDLOEPT 68  
 DB 126 SGGGNNAPARSPSPSTQPMWEHVNATQEARLLNLSRDTAENNETVEVISEMFDLOEPT 185  
 QY 69 CLOTRLELYKQGLRGSITLKLKGLPTWASHYKHQCPTPTTSCATQIITPESKFNKDF 128  
 DB 186 CLOTRLELYKQGLRGSITLKLKGLPTWASHYKHQCPTPTTSCATQIITPESKFNKDF 245  
 QY 129 LLVIPDCWEPVQES 142  
 DB 246 LLVIPDCWEPVQES 259  
 RESULT 5  
 ADJ14353  
 ID ADJ14353 standard; protein; 259 AA.  
 XX  
 AC ADJ14353;  
 XX

XX 20-MAY-2004 (first entry)  
DT Chimera protein related to human interleukin-3 mutant protein SEQ ID 141.  
XX stem cell; antianaemic; immunostimulant; immunomodulator;  
XX antiinflammatory; dermatological; immunosuppressive; cytostatic;  
KW neuroprotective; haemopoietic disorder; gene therapy; myeloid; erythroid;  
KW lymphoid; megakaryocyte; aplastic anaemia; periodic neutropenia;  
KW Chediak-Higashi syndrome; systemic lupus erythematosus; leukaemia;  
KW myelodysplastic syndrome; myelofibrosis; interleukin-3; IL-3; chimera.  
XX Unidentified.  
XX US2003185790-A1.  
XX 02-OCT-2003.  
XX 26-FEB-2002; 2002US-00083446.  
XX 24-NOV-1992; 92US-00981044.  
PR 22-NOV-1993; 93WO-US011197.  
PR 04-FEB-1994; 94US-00192325.  
PR 02-FEB-1995; 95WO-US001185.  
PR 06-APR-1995; 95US-00411795.  
PR 06-JUN-1995; 95US-00446872.  
PR 09-DEC-1996; 96US-00762227.  
XX (BAUE//) BAUER S C.  
PA (ABRA//) ABRAMS M A.  
PA (BRAF//) BRAFORD-GOLDBERG S R.  
PA (CAPA//) CAPARON M H.  
PA (EAST//) EASTON A M.  
PA (KLEI//) KLEIN B K.  
PA (MCKE//) MCKEARN J P.  
PA (OLIN//) OLINS P O.  
PA (PAIK//) PAIK K.  
PA (THOM//) THOMAS J W.  
XX  
XX Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
PI Klein BK, Mckearn JP, Olins PO, Paik K, Thomas JW;  
XX WPI; 2004-096775/10.  
XX  
XX Ex vivo expansion of stem cells, e.g. hematopoietic cells for treating  
PT aplastic anemia, involves culturing the stem cells with growth medium  
PT comprising chimera protein, and harvesting the cultured stem cells.  
XX  
XX Claim 8; SEQ ID NO 141; 202pp; English.  
XX  
XX The invention relates to a novel method whereby stem cells are ex vivo  
CC expanded via culturing the stem cells with a growth medium comprising a  
CC chimera protein, followed by harvesting of the cultured stem cells. The  
CC method of the invention has antianaemic, immunostimulant,  
CC immunomodulator, antiinflammatory, dermatological, immunosuppressive,  
CC cytostatic and neuroprotective applications and may be useful to target  
CC haemopoietic cells for gene therapy, preferably for treating patients  
CC having a haemopoietic disorder characterised by decreased levels of  
CC myeloid, erythroid, lymphoid, and/or megakaryocyte cells of haemopoietic  
CC system. The expanded ex vivo cells may be used to treat neutropenia,  
CC aplastic anaemia, periodic neutropenia, Chediak-Higashi syndrome,  
CC systemic lupus erythematosus, leukaemia, myelodysplastic syndrome or  
CC myelofibrosis. The current sequence is that of a chimera protein related  
CC to the human interleukin-3 (IL-3) mutant protein of the invention.  
XX  
XX Sequence 259 AA;  
SQ  
Query Match 89.2%; Score 682; DB 8; Length 259;  
Best Local Similarity 97.0%; Pred. No. 2.2e-64;  
Matches 130; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
9 SGIEGRMAPRSPSPSTQPEWHVNAIQEARRLLNLSRDTAAEMNETVEVISEMFDLQBP 68  
||| |||||||||||||||||||||||||||||||||||||||||||||||||||||||||

Db 126 SGGGNNMAPRSPSPSTQPEWHVNAIQEARRLLNLSRDTAAEMNETVEVISEMFDLQBP 185  
Qy 69 CLOTRLELYKQGLRGLTKLKGPLTMMASHYKQHCPPTPETSCATQIITFFSFKENLKDF 128  
Db 186 CLOTRLELYKQGLRGLTKLKGPLTMMASHYKQHCPPTPETSCATQIITFFSFKENLKDF 245  
Qy 129 LLVIPPDCWEPVQE 142  
Db 246 LLVIPPDCWEPVQE 259  
RESULT 6  
AAW19763  
ID AAW19763 standard; protein; 533 AA.  
XX  
AC AAW19763;  
XX  
DT 17-SEP-1997 (first entry)  
XX  
DE p53-GM-CSF immunostimulant fusion protein.  
XX  
KW p53-GM-CSF; granulocyte macrophage colony stimulating factor;  
KW tumour suppressor gene; immunostimulant; cancer; therapy.  
XX  
XX Homo sapiens.  
FH Key Location/Qualifiers  
FT Protein 1..393  
FT Peptide /label= p53  
FT /label= Linker 394..395  
FT /notes= "product of XbaI linker" /label= 396..524  
FT Protein 396..524  
FT /label= GM-CSF 525..533  
FT Peptide /note= "hexahistidine tag"  
XX  
PN WO9724438-A1.  
XX  
PD 10-JUL-1997.  
XX  
PF 23-DEC-1996; 96WO-US020241.  
XX  
PR 28-DEC-1995; 95US-00579823.  
XX  
PA (ACTI-) ACTIVATED CELL THERAPY INC.  
XX  
PI Laus R, Ruegg CL, Wu H;  
XX  
XX WPI; 1997-363674/33.  
DR N-PSDB; AAT72724.  
XX  
XX Potent APC that activates T-cells to give multivalent cellular immune  
PT response - can also induce a cytotoxic T-cell response in a vertebrate  
PT subject.  
XX  
XX Example 7; Fig 11; 45pp; English.  
XX  
CC A fusion protein (AAW19763) comprises human p53 tumour suppressor protein  
CC and granulocyte-macrophage colony stimulating factor (GM-CSF). It is the  
CC expression product of a nucleic acid molecule (AAT72724) prep'd. by PCR  
CC amplification of p53 cDNA GM-CSF cDNA sequences (the GM-CSF antisense  
CC primer including a hexahistidine tag sequence) and their fusion via a  
CC XbaI linker. Fusion expression vectors can be used to transfect mammalian  
CC and insect cells. The p53-GM-CSF fusion protein is used to generate anti-  
CC p53 immunity. Tumour cells are eliminated by cytotoxic T lymphocytes  
CC activated in vivo or in vitro by exposure to antigen- presenting cells  
CC exposed to the fusion protein  
XX  
SQ Sequence 533 AA;  
Query Match 88.8%; Score 679; DB 2; Length 533;  
Best Local Similarity 97.7%; Pred. No. 1.3e-63;

```
Matches 128; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
QY 12 EGMAPARSPSTQPEWHVNAIQEARRLLNLSRDAAENNETVEISEMFDLQPTCLQ 71
DB : |||||
393 DRSAPARSPSTQPEWHVNAIQEARRLLNLSRDAAENNETVEISEMFDLQPTCLQ 452
QY 72 TRLELYKQGLRGLTKLKGPLTWASHYKQHCPTTSCATQIITFESFKNLKDPLLV 131
DB |||||
453 TRLELYKQGLRGLTKLKGPLTWASHYKQHCPTTSCATQIITFESFKNLKDPLLV 512
QY 132 IPFDCWEPVQE 142
DB |||||
513 IPFDCWEPVQE 523

RESULT 7
AAP90118
ID AAP90118 standard; protein; 128 AA.
XX
AC AAP90118;
XX
XX 25-MAR-2003 (revised)
DT 01-NOV-1989 (first entry)
XX
XX Synthetic human granulocyte colony stimulating factor.
XX
XX BspMI; restriction sites; blunt ends; fusion proteins; synthetic;
KW human granulocyte colony stimulating factor.
XX
XX Homo sapiens.
OS Synthetic.
XX
XX GB2212160-A.
PN
XX 19-JUL-1989.
PD
XX 13-NOV-1987; 87GB-00726581.
PF
XX 13-NOV-1987; 87GB-00026581.
PR
XX (BRBI-) BRITISH BIO-TECHN L.
PA
XX Edwards RM;
PI
XX
XX WPI; 1989-208959/29.
DR
XX N-PSDB; AAN90383.
XX
XX DNA including recognition site for BspMI enzyme - allowing generation of
PT blunt end for fusion in prodn. of fusion proteins.
PT
XX
XX Disclosure; Fig 2; 23pp; English.
PS
XX Synthetic human granulocyte colony stimulating factor (GM-CSF) contg.
CC useful restriction sites, and a BspMI site. See corresp. AAN90383. Its
CC DNA is useful for generating blunt ends in fusion protein prodn. (Updated
CC on 25-MAR-2003 to correct PF field.) (Updated on 25-MAR-2003 to correct
CC PR field.) (Updated on 25-MAR-2003 to correct PA field.)
XX
XX Sequence 128 AA;
SQ

Query Match 88.6%; Score 678; DB 1; Length 128;
Best Local Similarity 100.0%; Pred. No. 2.3e-64;
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 15 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAENNETVEISEMFDLQPTCLQ 74
DB 1 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAENNETVEISEMFDLQPTCLQ 60
QY 75 ELYKQGLRGLTKLKGPLTWASHYKQHCPTTSCATQIITFESFKNLKDPLLV 134
DB 61 ELYKQGLRGLTKLKGPLTWASHYKQHCPTTSCATQIITFESFKNLKDPLLV 120
QY 135 DCWEPVQE 142
DB 121 DCWEPVQE 128

RESULT 9
AAR79338
ID AAR79338 standard; protein; 128 AA.
XX
AC AAR79338;
XX
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|||||
DB 121 DCWEPVQE 128

RESULT 8
AAP90115
ID AAP90115 standard; protein; 128 AA.
XX
AC AAP90115;
XX
XX 25-MAR-2003 (revised)
DT 01-NOV-1989 (first entry)
XX
XX Synthetic human granulocyte colony stimulating factor.
XX
XX Human granulocyte colony stimulating factor; Synthetic restriction sites;
KW cassette mutagenesis; GM-CSF; expression system.
XX
XX Homo sapiens.
OS Synthetic.
XX
XX GB2212159-A.
PN
XX 19-JUL-1989.
PD
XX 13-NOV-1987; 87GB-00026580.
PF
XX 13-NOV-1987; 87GB-00026580.
PR
XX (BRBI-) BRITISH BIO-TECHN L.
PA
XX Edwards RM;
PI
XX
XX WPI; 1989-208958/29.
DR
XX Human granulocyte-macrophage colony stimulating factor - synthetic DNA
PT includes restriction sites for cassette mutagenesis and incorporation in
PT expression systems.
XX
XX Claim 2; Fig 3a; 21pp; English.
PS
XX Synthetic human granulocyte colony stimulating factor (GM-CSF), see
CC corresp. AAN90274. Its DNA has useful restriction sites for: HindIII;
CC BspMI; NcoI; BstEII; BsmI; EcoRV; BglI; ApaI; BclI; XbaI; BamHI; and
CC EcoRI. Used to facilitate cassette mutagenesis of selected regions.
CC Synthesised by phosphoramidite chemistry, by dividing desired gene into
CC 18 oligomers. (Updated on 25-MAR-2003 to correct PA field.)
XX
XX Sequence 128 AA;
SQ

Query Match 88.6%; Score 678; DB 1; Length 128;
Best Local Similarity 100.0%; Pred. No. 2.3e-64;
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 15 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAENNETVEISEMFDLQPTCLQ 74
DB 1 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAENNETVEISEMFDLQPTCLQ 60
QY 75 ELYKQGLRGLTKLKGPLTWASHYKQHCPTTSCATQIITFESFKNLKDPLLV 134
DB 61 ELYKQGLRGLTKLKGPLTWASHYKQHCPTTSCATQIITFESFKNLKDPLLV 120
QY 135 DCWEPVQE 142
DB 121 DCWEPVQE 128

RESULT 9
AAR79338
ID AAR79338 standard; protein; 128 AA.
XX
AC AAR79338;
XX
```

DT 25-AUG-1999 (first entry)  
XX  
DE PMON13012 peptide.  
XX  
KW Interleukin; hIL-3; CSF; colony stimulating factor; cytokine; lymphokine;  
KW mutant; mutein; fusion protein.  
XX  
OS Synthetic.  
OS Homo sapiens.  
XX  
PN W05951254-A1.  
XX  
XX 10-AUG-1995.  
XX  
XX 02-FEB-1995; 95NO-US001185.  
XX  
XX 04-FEB-1994; 94US-00192325.  
XX  
XX (SEAR ) SEARLE & CO G D.  
XX  
XX Bauer CS, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
PI Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;  
PI  
XX WPI; 1995-283774/37.  
DR N-PSDB; AAQ97208.  
XX  
XX Fusion proteins comprising a human interleukin-3 variant, a linker and  
PT interleukin-3, a variant or a colony stimulating factor - useful to  
PT increase haematopoietic cell prodn. in a mammal.  
XX  
XX Example 20; Page 125; 447pp; English.  
PS  
XX A new fusion protein is disclosed which has the formula R1-L-R2, R2-L-R1,  
CC R1-R2, R2-R1, R1-L-R1 or R1-R1, where R1 is a mutant or variant of human  
CC interleukin-3 (hIL-3), R2 is a second colony stimulating factor (CSF)  
CC including cytokine, lymphokine, interleukin, haematopoietic growth factor  
CC or IL-3 variant, and L is a linker. Generic sequences are described in  
CC AA003235 - AA003242, and specifically claimed examples are shown in  
CC AA079298-R79335 and AA079342-R79345. The fusion protein is made by  
CC recombinant DNA techniques. Specifically claimed examples of DNA  
CC sequences which encode these proteins are shown in AAQ97167-Q97204 and  
CC AAQ97222-Q97227. The fusion protein is used to increase haematopoietic  
CC cell production. It is also useful as an IL-3 antagonist or as a discrete  
CC antigenic fragment for production of antibodies useful in immunoassays  
CC and immunotherapy. Antagonists are used to block the growth of certain  
CC cancer cells and in treatment of asthma. The fusion protein can also be  
CC used to stimulate bone marrow and blood cell activation and growth in  
CC vitro before infusion; and to treat diseases characterised by decreased  
CC levels of myeloid, erythroid, lymphoid and/or megakaryocyte cells of the  
CC haematopoietic system. The protein has the usual activity of both its  
CC component proteins, but may have increased synergistic activity and  
CC reduced undesired side effects  
XX  
SQ Sequence 128 AA;  
Query Match 88.6%; Score 678; DB 2; Length 128;  
Best Local Similarity 100.0%; Pred. No. 2.3e-64;  
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 15 MAPARSPSPSTQPWVHNVAIQEARRLLNLSRDTAAEMNETVEISEMFDLQEPCTCQLTRL 74  
DB 1 MAPARSPSPSTQPWVHNVAIQEARRLLNLSRDTAAEMNETVEISEMFDLQEPCTCQLTRL 60  
QY 75 ELYKQGLRGLSTKLKGLPTWASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 134  
DB 61 ELYKQGLRGLSTKLKGLPTWASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 120  
QY 135 DCWEPVQE 142  
DB 121 DCWEPVQE 128  
RESULT 11  
ID AA553217  
ID AA553217 standard; protein; 128 AA.  
XX  
XX AA553217;  
XX AC  
XX DT 19-MAY-2000 (first entry)  
XX Human G-CSF mutant protein sequence SEQ ID NO:160.  
XX Human; interleukin 3; IL-3; mutant; mutein; CSF; cytokine;  
KW colony stimulating factor; haematopoietic growth factor; lymphokine;  
KW fusion protein; haematopoietic disorder; infection; cancer;

AAW00103  
ID AAW00103 standard; protein; 128 AA.  
XX  
AC AAW00103;  
XX  
DT 25-MAR-2003 (revised)  
DT 11-FEB-1997 (first entry)  
XX  
XX Granulocyte macrophage colony-stimulating factor (ile101).  
XX GM-CSF; granulocyte macrophage colony-stimulating factor; expression;  
KW construct; stable; production.  
KW  
XX Homo sapiens.  
XX JP08173185-A.  
XX  
PD 09-JUL-1996.  
XX  
XX 28-APR-1987; 95JP-00263370.  
XX  
XX 28-APR-1987; 87JP-00106148.  
XX  
XX (AMGE-) AMGEN INC.  
PA (KIRI ) KIRIN BREWERY KK.  
PA  
XX WPI; 1996-365600/37.  
DR N-PSDB; AAT34400.  
XX  
XX Prodn. of human granulocyte macrophage colony-stimulating factor - by  
PT culturing E. coli transformed with human GM-CSF DNA.  
PT  
XX Claim 1; Page 2; 16pp; Japanese.  
PS  
XX The present sequence is that of human granulocyte macrophage colony-  
CC stimulating factor (hGM-CSF; n = 101, Ile). A series of oligonucleotides  
CC were synthesised and ligated together to form a stable expression  
CC construct. The technique is used for the efficient prodn. of a  
CC glycoprotein with hGM-CSF activity. (Updated on 25-MAR-2003 to correct PF  
CC field.)  
XX  
SQ Sequence 128 AA;  
Query Match 88.6%; Score 678; DB 2; Length 128;  
Best Local Similarity 100.0%; Pred. No. 2.3e-64;  
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 15 MAPARSPSPSTQPWVHNVAIQEARRLLNLSRDTAAEMNETVEISEMFDLQEPCTCQLTRL 74  
DB 1 MAPARSPSPSTQPWVHNVAIQEARRLLNLSRDTAAEMNETVEISEMFDLQEPCTCQLTRL 60  
QY 75 ELYKQGLRGLSTKLKGLPTWASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 134  
DB 61 ELYKQGLRGLSTKLKGLPTWASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 120  
QY 135 DCWEPVQE 142  
DB 121 DCWEPVQE 128  
RESULT 11  
ID AA553217  
ID AA553217 standard; protein; 128 AA.  
XX  
XX AA553217;  
XX AC  
XX DT 19-MAY-2000 (first entry)  
XX Human G-CSF mutant protein sequence SEQ ID NO:160.  
XX Human; interleukin 3; IL-3; mutant; mutein; CSF; cytokine;  
KW colony stimulating factor; haematopoietic growth factor; lymphokine;  
KW fusion protein; haematopoietic disorder; infection; cancer;

KW radiation therapy; chemotherapy; bone marrow suppressive drug;  
 KW bone marrow activation; blood cell activation; blood transplant.

XX Homo sapiens.  
 OS Synthetic.

XX US6022535-A.

XX 08-FEB-2000.

XX 06-JUN-1995; 95US-00469318.

XX 04-FEB-1994; 94US-00192325.

PR 02-FEB-1995; 95WO-US001185.

PR 06-APR-1995; 95US-00411795.

XX (SEAR ) SEARLE & CO G D.

XX Bauer SC, Abrams MA, Brafard-Goldberg SR, Easton AM, Klein BK;

PI Paik K, Thomas JW, Mckearn JP, Olins PO, Caparon MH;

XX WPI; 2000-160368/14.

XX Treating hematopoietic disorders with fusion proteins comprising mutated  
 PT interleukin-3 fused with secondary colony stimulating factors or other  
 PT interleukin-3 variants.

XX Example 20; Col 109-111; 276pp; English.

XX Methods have been developed for treating haematopoietic disorders with  
 CC fusion proteins comprising recombinant, mutated human interleukin-3 (IL-  
 CC 3) variants or mutant proteins (mutins) fused with secondary colony  
 CC stimulating factors (CSFs) (e.g. cytokines, lymphokines, interleukin  
 CC and/or haematopoietic colony stimulating factors) or other interleukin-3  
 CC variants with or without a linker. The methods may be used in vivo to  
 CC treat haematopoietic disorders resulting from bacterial, viral and fungal  
 CC infections, cancer radiation therapy, chemotherapy or bone marrow  
 CC suppressive drugs. They may also be used in vitro to stimulate bone  
 CC marrow and blood cell activation and growth prior to infusion of the bone  
 CC marrow and blood transplants into patients. IL-3 is a haematopoietic  
 CC growth factor which has the property of being able to promote the  
 CC survival, growth and differentiation of haematopoietic cells. The fusion  
 CC molecules are characterised by possessing the usual activity of both of  
 CC their constituent peptides and further by having a biological or  
 CC physiological activity greater than the additive function of the IL-3 or  
 CC second CSF alone (i.e. the peptides act synergistically). Their activity  
 CC may also be further enhanced by the mutations they comprise. The  
 CC variations may further reduce undesirable side effects associated with IL  
 CC -3. AAY53130 to AAY53226, and AAA03721 to AAA03782 represent sequences  
 CC used in the exemplification of the present invention

XX Sequence 128 AA;

Query Match 88.6%; Score 678; DB 3; Length 128;  
 Best Local Similarity 100.0%; Pred. No. 2.3e-64;  
 Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPSTQWEHVNAIQEARLLNLSRDTAAENNETVEVISEMFDLQETCLQTRL 74

DB 1 MAPARSPSPSTQWEHVNAIQEARLLNLSRDTAAENNETVEVISEMFDLQETCLQTRL 60

QY 75 ELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 134

DB 61 ELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 120

QY 135 DCWEPVQE 142

DB 121 DCWEPVQE 128

RESULT 12

AAE14011

ID AAE14011 standard; protein; 128 AA.

XX  
 AC

XX AAE14011;

XX 26-FEB-2002 (first entry)

XX Chemically modified myelopoietin (MPO) conjugate related protein #11.

XX Myelopoietin conjugate; MPO; immunosuppressive; vulnery; antiparasitic;  
 KW antibacterial; virucide; interleukin-3; IL-3; haematopoietic disorder;  
 KW haematopoietic growth factor receptor; neutropenia; thrombocytopenia;  
 KW leukopenia; anaemia; chemotherapy; bone marrow transplantation; burn;  
 KW radiation therapy; haematopoietic progenitor mobilisation; infection;  
 KW wound healing.

XX Unidentified.

XX WO200176639-A2.

XX 18-OCT-2001.

XX 06-APR-2001; 2001WO-US011256.

XX 06-APR-2000; 2000US-0195496P.

XX (PHAA ) PHARMACIA CORP.

XX (FINN/) FINN R.

XX (GOKA/) GOKARN Y.

XX (HILL/) HILLS R.

XX (NICAR/) NICASTRO P.

XX (QIHH/) QI H.

XX (SEDO/) SEDO K.

XX (SIEG/) SIEGEL N.

XX (WALT/) WALTER S.

XX Finn R, Gokarn Y, Hills R, Nicastro P, Qi H, Sedo K, Siegel N;

XX Walter S;

XX WPI; 2001-657130/75.

XX Myelopoietin conjugate for treating e.g. leukopenia comprises a water-  
 PT soluble polymer attached to the protein.

XX Disclosure; Page 215; 429pp; English.

XX The invention relates to chemically modified myelopoietin (MPO)  
 CC conjugates having at least one water-soluble polymer molecule covalently  
 CC attached, via activating group, to at least one amino acid residue of a  
 CC biologically active myelopoietin polypeptide. MPOs are multifunctional  
 CC agonists of human interleukin-3 (IL-3) and another haematopoietic growth  
 CC factor receptor. Sequences of the invention are useful for treating  
 CC haematopoietic disorders (e.g. neutropenia, leukopenia, thrombocytopenia  
 CC and anaemia), including those arising from chemotherapy and radiation  
 CC therapy. MPOs are also useful in bone marrow transplantation, wound  
 CC healing, burn treatment and the treatment of parasite, bacterial or viral  
 CC infection. They are useful for mobilising haematopoietic progenitors and  
 CC stem cells. The chemically modified MPOs have a longer lasting neutrophil  
 CC releasing activity, decreased clearance rate, increased stability and  
 CC decreased antigenicity than unmodified myelopoietins. The present  
 CC sequence is a chemically modified myelopoietin conjugate related protein

XX Sequence 128 AA;

Query Match 88.6%; Score 678; DB 4; Length 128;  
 Best Local Similarity 100.0%; Pred. No. 2.3e-64;  
 Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPSTQWEHVNAIQEARLLNLSRDTAAENNETVEVISEMFDLQETCLQTRL 74

DB 1 MAPARSPSPSTQWEHVNAIQEARLLNLSRDTAAENNETVEVISEMFDLQETCLQTRL 60

QY 75 ELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 134

DB 61 ELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 120

QY 135 DCWEPVQE 142  
 Db 121 DCWEPVQE 128

RESULT 13  
 ABG97784  
 ID ABG97784 standard; protein; 128 AA.  
 AC ABG97784;  
 XX  
 XX  
 XX 18-DEC-2002 (first entry)  
 XX Human Interleukin-3 chimaeric protein #39.  
 DE Human; interleukin-3; IL-3; mutant; mutein; stem cell;  
 KW haematopoietic factor; GM-CSF; colony stimulating factor; CSF-1; G-CSF;  
 KW G-CSFser17; c-mpl ligand; TPO; MGDF; erythropoietin; flt3 ligand;  
 KW human growth hormone; B-cell growth factor; leukaemia;  
 KW B-cell differentiation factor; eosinophil differentiation factor;  
 KW stem cell factor; SCF; cyclic neutropenia; aplastic anaemia;  
 KW thrombocytopenia; idiopathic neutropenia; Chediak-Higashi syndrome;  
 KW systemic lupus erythematosus; SLE; myelodysplastic syndrome;  
 KW myelofibrosis.  
 XX  
 XX Homo sapiens.  
 OS Synthetic.  
 OS  
 XX US6436387-B1.  
 XX  
 XX 20-AUG-2002.  
 PD  
 XX 09-DEC-1996; 96US-00762227.  
 XX  
 XX 24-NOV-1992; 92US-00981044.  
 PR 22-NOV-1993; 93WO-US011197.  
 PR 04-FEB-1994; 94US-00192325.  
 PR 04-FEB-1995; 95WO-US001185.  
 PR 06-APR-1995; 95US-00411795.  
 PR 06-JUN-1995; 95US-00446872.  
 XX (SEAR ) SEARLE & CO G D.  
 XX  
 XX Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
 PI Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;  
 XX WPI; 2002-749206/81.  
 XX  
 XX Ex vivo expansion of stem cells, for enhancing transduction efficiency of  
 PT cultured stem cells, comprises culturing stem cells in growth medium  
 PT having mutant interleukin-3, and hematopoietic factor, and harvesting  
 PT cultured cells.  
 XX  
 XX Claim 8; Col 283-284; 203pp; English.  
 PS  
 XX The invention relates to ex vivo expansion of stem cells, comprises  
 CC culturing stem cells with a growth medium comprising a chimaera protein,  
 CC and harvesting the cultured stem cells. The chimaera is based on a  
 CC mutated human interleukin-3 (IL-3) sequence coupled to a haematopoietic  
 CC factor (e.g. GM-CSF (colony stimulating factor), CSF-1, G-CSF, G-  
 CC CSFser17, c-mpl ligand TPO, MGDF, erythropoietin, IL-1-13, IL-15, IL-16,  
 CC flt3 ligand, human growth hormone, B-cell growth factor, B-cell  
 CC differentiation factor, eosinophil differentiation factor and stem cell  
 CC factor (SCF)) via a peptide linker. The formula for the chimaera is given  
 CC in the specification. Also included is a method for enhancing the  
 CC efficiency of the transduction of cultured stem cells by a heterologous  
 CC gene, comprising: (a) removing stem cells from a patient or donor; (b)  
 CC culturing the stem cells with a growth medium comprising the chimaera (c)  
 CC transducing DNA into cultured cells; and (d) harvesting the transduced  
 CC cells. The method is useful for ex vivo expansion of stem cells, and  
 CC enhancing the efficiency of the transduction of cultured stem cells by a  
 CC heterologous gene. The method is also useful for treating a patient

CC having a haematopoietic disorder. The expanded haematopoietic cells are  
 CC also useful in the treatment of cyclic neutropenia, aplastic anaemia,  
 CC thrombocytopenia, idiopathic neutropenia, Chediak-Higashi syndrome,  
 CC systemic lupus erythematosus (SLE), leukaemia, myelodysplastic syndrome  
 CC and myelofibrosis. The present sequence is a human IL-3  
 CC mutant/haematopoietic factor chimaeric sequence  
 XX  
 XX Sequence 128 AA;  
 SQ  
 Query Match 88.6%; Score 678; DB 5; Length 128;  
 Best Local Similarity 100.0%; Pred. No. 2.3e-64;  
 Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 15 MAPARSPSPSTQWEHVNAIQEARRLLNLSRDTAAEMNETVEVISEMFDLQPTCLQTRL 74  
 Db 1 MAPARSPSPSTQWEHVNAIQEARRLLNLSRDTAAEMNETVEVISEMFDLQPTCLQTRL 60  
 QY 75 ELYKQGLRSLTKLKGPLTMASHYKQHCPPPTSCATQITTFESFKENLKDFLVIPP 134  
 Db 61 ELYKQGLRSLTKLKGPLTMASHYKQHCPPPTSCATQITTFESFKENLKDFLVIPP 120  
 QY 135 DCWEPVQE 142  
 Db 121 DCWEPVQE 128

RESULT 14  
 ADJ14372  
 ID ADJ14372 standard; protein; 128 AA.  
 XX  
 AC ADJ14372;  
 XX  
 XX 20-MAY-2004 (first entry)  
 DT  
 DE Protein related to human interleukin-3 (IL-3) mutant protein SEQ ID 140.  
 XX  
 XX stem cell; antianaemic; immunostimulant; immunomodulator;  
 KW antiinflammatory; dermatological; immunosuppressive; cytostatic;  
 KW neuroprotective; haematopoietic disorder; gene therapy; myeloid; erythroid;  
 KW lymphoid; megakaryocyte; aplastic anaemia; periodic neutropenia;  
 KW Chediak-Higashi syndrome; systemic lupus erythematosus; leukaemia;  
 KW myelodysplastic syndrome; myelofibrosis; interleukin-3; IL-3.  
 XX  
 OS Unidentified.  
 XX  
 XX US2003185790-A1.  
 XX  
 XX 02-OCT-2003.  
 PD  
 XX 26-FEB-2002; 2002US-00083446.  
 XX  
 XX 24-NOV-1992; 92US-00981044.  
 PR 22-NOV-1993; 93WO-US011197.  
 PR 04-FEB-1994; 94US-00192325.  
 PR 02-FEB-1995; 95WO-US001185.  
 PR 06-APR-1995; 95US-00411795.  
 PR 06-JUN-1995; 95US-00446872.  
 PR 09-DEC-1996; 96US-00762227.  
 XX (BAUE/) BAUER S C.  
 PA (ABRA/) ABRAMS M A.  
 PA (BRAP/) BRAFORD-GOLDBERG S R.  
 PA (CAPA/) CAPARON M H.  
 PA (EAST/) EASTON A M.  
 PA (KLEI/) KLEIN B K.  
 PA (MCKE/) MCKEARN J P.  
 PA (OLIN/) OLINS P O.  
 PA (PAIK/) PAIK K.  
 PA (THOM/) THOMAS J W.  
 XX  
 XX Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
 PI Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;  
 XX



DR WPI; 2004-096775/10.

XX Ex vivo expansion of stem cells, e.g. hematopoietic cells for treating

PT aplastic anemia, involves culturing the stem cells with growth medium

PT comprising chimera protein, and harvesting the cultured stem cells.

XX

PS Disclosure; SEQ ID NO 160; 202pp; English.

XX

CC The invention relates to a novel method whereby stem cells are ex vivo

CC expanded via culturing the stem cells with a growth medium comprising a

CC chimera protein, followed by harvesting of the cultured stem cells. The

CC method of the invention has antianaemic, immunostimulant,

CC immunomodulator, antiinflammatory, dermatological, immunosuppressive,

CC cytostatic and neuroprotective applications and may be useful to target

CC hematopoietic cells for gene therapy, preferably for treating patients

CC having a haemopoietic disorder characterised by decreased levels of

CC myeloid, erythroid, lymphoid, and/or megakaryocyte cells of haemopoietic

CC system. The expanded ex vivo cells may be used to treat neutropenia,

CC aplastic anaemia, periodic neutropenia, Chediak-Higashi syndrome,

CC systemic lupus erythematosus, leukaemia, myelodysplastic syndrome or

CC myelofibrosis. The current sequence is that of a protein related to the

CC human interleukin-3 (IL-3) mutant protein of the invention.

XX

SQ Sequence 128 AA;

Query Match 88.6%; Score 678; DB 8; Length 128;

Best Local Similarity 100.0%; Pred. No. 2.3e-64;

Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAMNETVEVISEMFDLQETCLOTRL 74

DB 1 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAMNETVEVISEMFDLQETCLOTRL 60

QY 75 ELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 134

DB 61 ELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 120

QY 135 DCWEPVQE 142

DB 121 DCWEPVQE 128

RESULT 15

AAR79320

ID AAR79320 standard; protein; 274 AA.

XX

AC AAR79320;

XX

DT 25-AUG-1999 (first entry)

XX

DE IL-3 containing fusion protein.

XX

KW interleukin; hIL-3; CSF; colony stimulating factor; cytokine; lymphokine;

KW mutant; mutein; fusion protein.

XX

OS Synthetic.

XX

PN WO9521254-A1.

XX

PD 10-AUG-1995.

XX

PF 02-FEB-1995; 95WO-US0001185.

XX

PR 04-FEB-1994; 94US-00192325.

XX

PA (SEAR ) SEARLE & CO G D.

XX

PI Bauer CS, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;

PI Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;

XX

DR WPI; 1995-283774/37.

DR N-PSDB; AAQ97183.

XX

PT Fusion proteins comprising a human interleukin-3 variant, a linker and

PT interleukin-3, a variant or a colony stimulating factor - useful to

XX increase haematopoietic cell prodn. in a mammal.

PS Claim 16; Page 87-88; 447pp; English.

XX

CC A new fusion protein has the formula R1-L-R2, R2-L-R1, R1-R2, R2-R1, R1-L

CC -R1 or R1-R1, in which R1 is a mutant or variant of human interleukin-3

CC (hIL-3) having the present generic sequence, R2 is a second colony

CC stimulating factor (CSF) including cytokine, lymphokine, interleukin,

CC haematopoietic growth factor or IL-3 variant, and L is a linker. The

CC present sequence corresponds to native hIL-3(1-133) in which 1-14 amino

CC acids may be deleted from the N-terminal, 1-15 amino acids can be deleted

CC from the C-terminal, and at least 4 and up to 44 amino acids in the

CC region 17-123 are different from those in the native protein. The fusion

CC protein is used to increase haematopoietic cell production. It is also

CC useful as an IL-3 antagonist or as a discrete antigenic fragment for

CC production of antibodies useful in immunoassays and immunotherapy.

CC Antagonists are used to block the growth of certain cancer cells and in

CC treatment of asthma. The fusion protein can also be used to stimulate

CC bone marrow and blood cell activation and growth in vitro before infusion

CC ; and to treat diseases characterised by decreased levels of myeloid,

CC erythroid, lymphoid and/or megakaryocyte cells of the haematopoietic

CC system. The protein has the usual activity of both its component

CC proteins, but may have increased synergistic activity and reduced

CC undesired side effects

XX

SQ Sequence 274 AA;

Query Match 88.6%; Score 678; DB 2; Length 274;

Best Local Similarity 100.0%; Pred. No. 6.4e-64;

Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAMNETVEVISEMFDLQETCLOTRL 74

DB 147 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAMNETVEVISEMFDLQETCLOTRL 206

QY 75 ELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 134

DB 207 ELYKQGLRGLSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 266

QY 135 DCWEPVQE 142

DB 267 DCWEPVQE 274

Search completed: March 8, 2005, 16:10:22

Job time : 167 secs



GenCore version 5.1.6  
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OM protein - protein search, using sw model

Run on: March 8, 2005, 16:13:34 ; Search time 131 Seconds  
(without alignments)  
356.675 Million cell updates/sec

Title: US-10-723-083-2  
Perfect score: 765  
Sequence: 1 MHHHHSGSGEGRMAPARS.....ENLKDFLLVLPDCWEPVQE 142

Scoring table:

BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1391452 seqs, 329044822 residues

Total number of hits satisfying chosen parameters: 1391452

Minimum DB seq length: 0

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Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA.\*  
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19: /cgn2\_6/ptodata/2/pubpaa/US60\_NEW\_PUB.pep.\*  
20: /cgn2\_6/ptodata/2/pubpaa/US60\_PUBCOMB.pep.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	682	89.2	259	14	US-10-083-446-141 Sequence 141, App
2	678	88.6	128	14	US-10-083-446-160 Sequence 160, App
3	678	88.6	274	14	US-10-083-446-144 Sequence 144, App
4	678	88.6	301	14	US-10-083-446-142 Sequence 142, App
5	675	88.2	712	15	US-10-609-346-10 Sequence 10, Appl
6	673	88.0	127	9	US-09-821-883-18 Sequence 18, Appl
7	673	88.0	127	9	US-09-800-016-1 Sequence 1, Appl
8	673	88.0	127	9	US-09-792-793A-15 Sequence 15, Appl
9	673	88.0	127	14	US-10-400-377-8 Sequence 8, Appl
10	673	88.0	127	14	US-10-400-708-8 Sequence 8, Appl
11	673	88.0	127	14	US-10-298-148-8 Sequence 8, Appl
12	673	88.0	127	15	US-10-375-209A-15 Sequence 15, Appl
13	673	88.0	127	16	US-10-658-834A-202 Sequence 202, App

14	673	88.0	127	16	US-10-743-295-5 Sequence 5, Appli
15	673	88.0	127	16	US-10-773-939-8 Sequence 8, Appli
16	673	88.0	127	16	US-10-774-149-8 Sequence 8, Appli
17	673	88.0	143	15	US-10-449-831A-142 Sequence 142, App
18	673	88.0	144	9	US-09-923-246-114 Sequence 114, App
19	673	88.0	144	14	US-10-295-723-114 Sequence 114, App
20	673	88.0	144	14	US-10-282-622-10 Sequence 10, Appl
21	673	88.0	144	14	US-10-131-985-15 Sequence 15, Appl
22	673	88.0	144	15	US-10-116-275-217 Sequence 217, App
23	673	88.0	144	15	US-10-456-780-10 Sequence 10, Appl
24	673	88.0	144	15	US-10-411-037-18 Sequence 18, Appl
25	673	88.0	144	15	US-10-609-346-20 Sequence 20, Appl
26	673	88.0	144	15	US-10-411-026-18 Sequence 18, Appl
27	673	88.0	144	15	US-10-447-315-19 Sequence 19, Appl
28	673	88.0	144	15	US-10-410-962-18 Sequence 18, Appl
29	673	88.0	144	15	US-10-411-049-18 Sequence 18, Appl
30	673	88.0	144	16	US-10-659-684-114 Sequence 114, App
31	673	88.0	144	16	US-10-410-930-18 Sequence 18, Appl
32	673	88.0	144	16	US-10-410-997-18 Sequence 18, Appl
33	673	88.0	144	16	US-10-411-042-18 Sequence 18, Appl
34	673	88.0	144	16	US-10-287-994-18 Sequence 18, Appl
35	673	88.0	144	16	US-10-659-295-27 Sequence 27, Appl
36	673	88.0	144	16	US-10-410-913-18 Sequence 18, Appl
37	673	88.0	144	16	US-10-666-122-3 Sequence 3, Appli
38	673	88.0	144	16	US-10-666-122-5 Sequence 5, Appli
39	673	88.0	144	17	US-10-901-417-15 Sequence 15, Appl
40	673	88.0	144	17	US-10-410-980-18 Sequence 18, Appl
41	673	88.0	191	15	US-10-449-831A-188 Sequence 188, App
42	673	88.0	610	9	US-09-783-708-1 Sequence 1, Appli
43	673	88.0	690	9	US-09-821-883-2 Sequence 2, Appli
44	671	87.7	127	16	US-10-658-834A-376 Sequence 376, App
45	670	87.6	127	16	US-10-658-834A-362 Sequence 362, App

ALIGNMENTS

RESULT 1

US-10-083-446-141  
; Sequence 141, Application US/10083446  
; Publication No. US20030185790A1  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; Bauer, S. C.  
; Braford-Goldberg, Sarah R.  
; Caparon, Mair H.  
; Easton, Alan M.  
; Klein, Barbara K.  
; McKearn, John P.  
; Olin, Peter O.  
; Paik, Kuman  
; Thomas, John W.

TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells  
Using Multivariant (IL-3) Hematopoiesis Chimera Proteins

NUMBER OF SEQUENCES: 197

CORRESPONDENCE ADDRESS:

ADDRESSEE: S. Christopher Bauer, Pharmacia Corporation  
Corporate Patent Dept., Mail Zone 04E  
STREET: 800 N. Lindbergh  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63167

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

CURRENT APPLICATION NUMBER: US/10/083,446

FILING DATE: 26-Feb-2002

CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/762,227  
FILING DATE: 09-DEC-1996  
APPLICATION NUMBER: US 08/192,325  
FILING DATE: 14-FEB-1994  
APPLICATION NUMBER: US 08/446,872  
FILING DATE: 06-JUN-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: S. Christopher Bauer  
REGISTRATION NUMBER: 42,305  
REFERENCE/DOCKET NUMBER: C-2790/6  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (636)737-6257  
TELEFAX: (636)737-5452  
INFORMATION FOR SEQ ID NO: 141:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 259 amino acids  
TYPE: amino acid  
STRANDEDNESS: <Unknown>  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
SEQUENCE DESCRIPTION: SEQ ID NO: 141:  
US-10-083-446-141

Query Match 89.2%; Score 682; DB 14; Length 259;  
Best Local Similarity 97.0%; Pred. No. 7.2e-64;  
Matches 130; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 9 SGIEGRMAPARSPSTQPMHVNIAQEARLLNLSRDTAAMNETVEVISEMFDIQEPT 68  
DB 126 SGGGNNAPARSPSTQPMHVNIAQEARLLNLSRDTAAMNETVEVISEMFDIQEPT 185

QY 69 CLOTRLEYKQGRSLTKLKGPLTMASHYKQCPPTPETSCTOIITFESKFNKDF 128  
DB 186 CLOTRLEYKQGRSLTKLKGPLTMASHYKQCPPTPETSCTOIITFESKFNKDF 245

QY 129 LLVIPDCWEPVQE 142  
DB 246 LLVIPDCWEPVQE 259

RESULT 2  
US-10-083-446-160  
Sequence 160, Application US/10083446  
Publication No. US20030185790A1  
GENERAL INFORMATION:  
APPLICANT: Abrams, Mark A.  
Bauer, S. C.  
Braford-Goldberg, Sarah R.  
Caparon, Mair H.  
Easton, Alan M.  
Klein, Barbara K.  
McKearn, John P.  
Olin, Peter O.  
Paik, Kuman  
Thomas, John W.  
TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells  
Using Multivariant (IL-3) Hematopoiesis Chimera Proteins  
NUMBER OF SEQUENCES: 197  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: S. Christopher Bauer, Pharmacia Corporation  
Corporate Patent Dept., Mail Zone 04E  
STREET: 800 N. Lindbergh  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63167  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/10/083,446

FILING DATE: 26-Feb-2002  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/762,227  
FILING DATE: 09-DEC-1996  
APPLICATION NUMBER: US 08/192,325  
FILING DATE: 14-FEB-1994  
APPLICATION NUMBER: US 08/446,872  
FILING DATE: 06-JUN-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: S. Christopher Bauer  
REGISTRATION NUMBER: 42,305  
REFERENCE/DOCKET NUMBER: C-2790/6  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (636)737-6257  
TELEFAX: (636)737-5452  
INFORMATION FOR SEQ ID NO: 160:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 128 amino acids  
TYPE: amino acid  
STRANDEDNESS: <Unknown>  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
SEQUENCE DESCRIPTION: SEQ ID NO: 160:  
US-10-083-446-160

Query Match 88.6%; Score 678; DB 14; Length 128;  
Best Local Similarity 100.0%; Pred. No. 7.5e-64;  
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSTQPMHVNIAQEARLLNLSRDTAAMNETVEVISEMFDIQEPTCLOTRL 74  
DB 1 MAPARSPSTQPMHVNIAQEARLLNLSRDTAAMNETVEVISEMFDIQEPTCLOTRL 60

QY 75 ELYKQGRSLTKLKGPLTMASHYKQCPPTPETSCTOIITFESKFNKDFLLVIPF 134  
DB 61 ELYKQGRSLTKLKGPLTMASHYKQCPPTPETSCTOIITFESKFNKDFLLVIPF 120

QY 135 DCWEPVQE 142  
DB 121 DCWEPVQE 128

RESULT 3  
US-10-083-446-144  
Sequence 144, Application US/10083446  
Publication No. US20030185790A1  
GENERAL INFORMATION:  
APPLICANT: Abrams, Mark A.  
Bauer, S. C.  
Braford-Goldberg, Sarah R.  
Caparon, Mair H.  
Easton, Alan M.  
Klein, Barbara K.  
McKearn, John P.  
Olin, Peter O.  
Paik, Kuman  
Thomas, John W.  
TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells  
Using Multivariant (IL-3) Hematopoiesis Chimera Proteins  
NUMBER OF SEQUENCES: 197  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: S. Christopher Bauer, Pharmacia Corporation  
Corporate Patent Dept., Mail Zone 04E  
STREET: 800 N. Lindbergh  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63167  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/10/083,446  
FILING DATE: 26-Feb-2002  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/762,227  
FILING DATE: 09-DEC-1996  
APPLICATION NUMBER: US 08/192,325  
FILING DATE: 14-FEB-1994  
APPLICATION NUMBER: US 08/446,872  
FILING DATE: 06-JUN-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: S. Christopher Bauer  
REGISTRATION NUMBER: 42,305  
REFERENCE/DOCKET NUMBER: C-2790/6  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (636)737-6257  
TELEFAX: (636)737-5452  
INFORMATION FOR SEQ ID NO: 144:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 274 amino acids  
TYPE: amino acid  
STRANDEDNESS: <Unknown>  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
SEQUENCE DESCRIPTION: SEQ ID NO: 144:  
US-10-083-446-144

Query Match: 88.6%; Score 678; DB 14; Length 274;  
Best Local Similarity 100.0%; Pred. No. 2.1e-63;  
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAENETVEISEMFDLQETCLQTRL 74  
DB 147 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAENETVEISEMFDLQETCLQTRL 206

QY 75 ELYKQGLRSLTKLKGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 134  
DB 207 ELYKQGLRSLTKLKGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 266

QY 135 DCWEPVQE 142  
DB 267 DCWEPVQE 274

RESULT 4  
US-10-083-446-142  
Sequence 142, Application US/10083446  
Publication No. US20030185790A1  
GENERAL INFORMATION:  
APPLICANT: Abrams, Mark A.  
Bauer, S. C.  
Braford-Goldberg, Sarah R.  
Caparon, Mairé H.  
Easton, Alan M.  
Klein, Barbara K.  
McKearn, John P.  
Ollins, Peter O.  
Paik, Kuman  
Thomas, John W.  
TITLE OF INVENTION: Methods of Ex-Vivo Expansion Of Hematopoietic Cells  
Using Multivariant (IL-3) Hematopoiesis Chimera Proteins  
NUMBER OF SEQUENCES: 197  
CORRESPONDENCE ADDRESS:  
ADDRESSER: S. Christopher Bauer, Pharmacia Corporation  
STREET: 800 N. Lindbergh  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63167  
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/10/083,446  
FILING DATE: 26-Feb-2002  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/762,227  
FILING DATE: 09-DEC-1996  
APPLICATION NUMBER: US 08/192,325  
FILING DATE: 14-FEB-1994  
APPLICATION NUMBER: US 08/446,872  
FILING DATE: 06-JUN-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: S. Christopher Bauer  
REGISTRATION NUMBER: 42,305  
REFERENCE/DOCKET NUMBER: C-2790/6  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (636)737-6257  
TELEFAX: (636)737-5452  
INFORMATION FOR SEQ ID NO: 142:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 301 amino acids  
TYPE: amino acid  
STRANDEDNESS: <Unknown>  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
SEQUENCE DESCRIPTION: SEQ ID NO: 142:  
US-10-083-446-142

Query Match: 88.6%; Score 678; DB 14; Length 301;  
Best Local Similarity 100.0%; Pred. No. 2.3e-63;  
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAENETVEISEMFDLQETCLQTRL 74  
DB 174 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAENETVEISEMFDLQETCLQTRL 233

QY 75 ELYKQGLRSLTKLKGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 134  
DB 234 ELYKQGLRSLTKLKGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDPLLVIPF 293

QY 135 DCWEPVQE 142  
DB 294 DCWEPVQE 301

RESULT 5  
US-10-609-346-10  
Sequence 10, Application US/10609346  
Publication No. US20040063635A1  
GENERAL INFORMATION:  
APPLICANT: Yu, Zailin  
APPLICANT: Fu, Yan  
TITLE OF INVENTION: RECOMBINANT HUMAN ALBUMIN FUSION PROTEINS WITH LONG-LASTING BIOLO  
TITLE OF INVENTION: EFFECTS  
FILE REFERENCE: ZYU-0603  
CURRENT APPLICATION NUMBER: US/10/609,346  
CURRENT FILING DATE: 2003-06-26  
PRIOR APPLICATION NUMBER: US 60/392,948  
PRIOR FILING DATE: 2002-07-01  
NUMBER OF SEQ ID NOS: 40  
SOFTWARE: PatentIn version 3.1  
SEQ ID NO 10  
LENGTH: 712  
TYPE: PRT  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: HSA-GMCSF  
US-10-609-346-10

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Query Match      88.2%; Score 675; DB 15; Length 712;
Best Local Similarity 99.2%; Pred. No. 1.5e-62;
Matches 127; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPSTQWEHVNAIQEARLLNLSRDTAAEMNETVEVISEMFDLOEPTCLOTRL 74
Db 585 LAPARSPSPSTQWEHVNAIQEARLLNLSRDTAAEMNETVEVISEMFDLOEPTCLOTRL 644
QY 75 ELYKQGLRGSLLTKLKGPLTMMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 134
Db 645 ELYKQGLRGSLLTKLKGPLTMMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 704
QY 135 DCWEPVQE 142
Db 705 DCWEPVQE 712

RESULT 6
US-09-821-883-18
; Sequence 18, Application US/09821883
; Patent No. US20020061310A1
; GENERAL INFORMATION:
; APPLICANT: Laus, Reiner
; APPLICANT: Vidovic, Danir
; APPLICANT: Graddis, Thomas
; TITLE OF INVENTION: Compositions and Methods for Dendritic
; FILE REFERENCE: 7636-0022.30
; CURRENT APPLICATION NUMBER: US/09/821,883
; CURRENT FILING DATE: 2001-03-30
; PRIOR APPLICATION NUMBER: US 60/193,504
; PRIOR FILING DATE: 2000-03-30
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 18
; LENGTH: 127
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-821-883-18

Query Match      88.0%; Score 673; DB 9; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQWEHVNAIQEARLLNLSRDTAAEMNETVEVISEMFDLOEPTCLOTRL 75
Db 1 APARSPSPSTQWEHVNAIQEARLLNLSRDTAAEMNETVEVISEMFDLOEPTCLOTRL 60
QY 76 LYKQGLRGSLLTKLKGPLTMMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 135
Db 61 LYKQGLRGSLLTKLKGPLTMMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 120
QY 136 CWEPVQE 142
Db 121 CWEPVQE 127

RESULT 7
US-09-800-016-1
; Sequence 1, Application US/09800016
; Patent No. US20020141970A1
; GENERAL INFORMATION:
; APPLICANT: Pettit, Dean
; APPLICANT: Jochheim, Claudia
; TITLE OF INVENTION: STABLE AQUEOUS SOLUTIONS OF GRANULOCYTE MACROPHAGE COLONY-STIM
; FILE REFERENCE: 3253
; CURRENT APPLICATION NUMBER: US/09/800,016
; CURRENT FILING DATE: 2001-03-05
; NUMBER OF SEQ ID NOS: 1
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1
; LENGTH: 127
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; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-800-016-1

Query Match      88.0%; Score 673; DB 9; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQWEHVNAIQEARLLNLSRDTAAEMNETVEVISEMFDLOEPTCLOTRL 75
Db 1 APARSPSPSTQWEHVNAIQEARLLNLSRDTAAEMNETVEVISEMFDLOEPTCLOTRL 60
QY 76 LYKQGLRGSLLTKLKGPLTMMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 135
Db 61 LYKQGLRGSLLTKLKGPLTMMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 120
QY 136 CWEPVQE 142
Db 121 CWEPVQE 127

RESULT 8
US-09-792-793A-15
; Sequence 15, Application US/09792793A
; Patent No. US20020168370A1
; GENERAL INFORMATION:
; APPLICANT: McDonald, John R.
; APPLICANT: Coggin, Philip
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR TREATING SECONDARY TISSUE DAMAGE AND
; FILE REFERENCE: 25020-601D
; CURRENT APPLICATION NUMBER: US/09/792,793A
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 93
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 15
; LENGTH: 127
; TYPE: PRT
; ORGANISM: homo sapien
; FEATURE:
; OTHER INFORMATION: Human Chemokine Polypeptide: GM-CSF
US-09-792-793A-15

Query Match      88.0%; Score 673; DB 9; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQWEHVNAIQEARLLNLSRDTAAEMNETVEVISEMFDLOEPTCLOTRL 75
Db 1 APARSPSPSTQWEHVNAIQEARLLNLSRDTAAEMNETVEVISEMFDLOEPTCLOTRL 60
QY 76 LYKQGLRGSLLTKLKGPLTMMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 135
Db 61 LYKQGLRGSLLTKLKGPLTMMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPF 120
QY 136 CWEPVQE 142
Db 121 CWEPVQE 127

RESULT 9
US-10-400-377-8
; Sequence 8, Application US/10400377
; Publication No. US20030162949A1
; GENERAL INFORMATION:
; APPLICANT: Cox III, George N
; APPLICANT: Bolder Biotechnology, Inc.
; TITLE OF INVENTION: Derivatives of Growth Hormone and Related Proteins
; FILE REFERENCE: 4152-1-PUS
; CURRENT APPLICATION NUMBER: US/10/400,377
; CURRENT FILING DATE: 2003-03-26
; PRIOR APPLICATION NUMBER: US/09/462,941
; PRIOR FILING DATE: 2000-01-14
```



```
; PRIOR APPLICATION NUMBER: 60/052,516
; PRIOR FILING DATE: 1997-07-14
; NUMBER OF SEQ ID NOS: 41
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 8
; LENGTH: 127
; TYPE: PR
; ORGANISM: Homo sapiens
US-10-400-377-8

Query Match      88.0%; Score 673; DB 14; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQPWEHVNAIQEARRLLNLSRDTAAENNETVEVISEMFDLQEPCTCLOTRLE 75
Db 1 APARSPSPSTQPWEHVNAIQEARRLLNLSRDTAAENNETVEVISEMFDLQEPCTCLOTRLE 60

QY 76 LYKQGLRGSITKLGPLTWMAHYKQHCPTTPTSCATQIITPESPKENLKDFLLVIPP 135
Db 61 LYKQGLRGSITKLGPLTWMAHYKQHCPTTPTSCATQIITPESPKENLKDFLLVIPP 120

QY 136 CWPVQOE 142
Db 121 CWPVQOE 127

RESULT 10
US-10-400-708-8
; Sequence 8, Application US/10400708
; Publication No. US2003016685A1
; GENERAL INFORMATION:
; APPLICANT: Cox III, George N
; TITLE OF INVENTION: Derivatives of Growth Hormone and Related Proteins
; FILE REFERENCE: 4152-1-PUS
; CURRENT APPLICATION NUMBER: US/10/400,708
; PRIOR FILING DATE: 2003-03-26
; PRIOR APPLICATION NUMBER: US/09/462,941
; PRIOR FILING DATE: 2000-01-14
; PRIOR APPLICATION NUMBER: 60/052,516
; PRIOR FILING DATE: 1997-07-14
; NUMBER OF SEQ ID NOS: 41
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 8
; LENGTH: 127
; TYPE: PR
; ORGANISM: Homo sapiens
US-10-400-708-8

Query Match      88.0%; Score 673; DB 14; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQPWEHVNAIQEARRLLNLSRDTAAENNETVEVISEMFDLQEPCTCLOTRLE 75
Db 1 APARSPSPSTQPWEHVNAIQEARRLLNLSRDTAAENNETVEVISEMFDLQEPCTCLOTRLE 60

QY 76 LYKQGLRGSITKLGPLTWMAHYKQHCPTTPTSCATQIITPESPKENLKDFLLVIPP 135
Db 61 LYKQGLRGSITKLGPLTWMAHYKQHCPTTPTSCATQIITPESPKENLKDFLLVIPP 120

QY 136 CWPVQOE 142
Db 121 CWPVQOE 127

RESULT 11
US-10-298-148-8
; Sequence 8, Application US/10298148
; Publication No. US20030171284A1
; GENERAL INFORMATION:
; APPLICANT: Cox III, George N
```

```
; APPLICANT: Bolder Biotechnology, Inc.
; TITLE OF INVENTION: Derivatives of Growth Hormone and Related Proteins
; FILE REFERENCE: 4152-1-PUS
; CURRENT APPLICATION NUMBER: US/10/298,148
; CURRENT FILING DATE: 2002-11-15
; PRIOR APPLICATION NUMBER: US/09/462,941
; PRIOR FILING DATE: 2000-01-14
; PRIOR APPLICATION NUMBER: 60/052,516
; PRIOR FILING DATE: 1997-07-14
; NUMBER OF SEQ ID NOS: 41
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 8
; LENGTH: 127
; TYPE: PR
; ORGANISM: Homo sapiens
US-10-298-148-8

Query Match      88.0%; Score 673; DB 14; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQPWEHVNAIQEARRLLNLSRDTAAENNETVEVISEMFDLQEPCTCLOTRLE 75
Db 1 APARSPSPSTQPWEHVNAIQEARRLLNLSRDTAAENNETVEVISEMFDLQEPCTCLOTRLE 60

QY 76 LYKQGLRGSITKLGPLTWMAHYKQHCPTTPTSCATQIITPESPKENLKDFLLVIPP 135
Db 61 LYKQGLRGSITKLGPLTWMAHYKQHCPTTPTSCATQIITPESPKENLKDFLLVIPP 120

QY 136 CWPVQOE 142
Db 121 CWPVQOE 127

RESULT 12
US-10-375-209A-15
; Sequence 15, Application US/10375209A
; Publication No. US20030215421A1
; GENERAL INFORMATION:
; APPLICANT: McDonald, John R.
; APPLICANT: Coggins, Philip
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR TREATING SECONDARY TISSUE DAMAGE AND
; TITLE OF INVENTION: OTHER INFLAMMATORY CONDITIONS AND DISORDERS
; FILE REFERENCE: 25020-601E
; CURRENT APPLICATION NUMBER: US/10/375,209A
; CURRENT FILING DATE: 2003-02-24
; NUMBER OF SEQ ID NOS: 93
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 15
; LENGTH: 127
; TYPE: PR
; ORGANISM: homo sapien
; FEATURE:
; OTHER INFORMATION: Human Chemokine Polypeptide: GM-CSF
US-10-375-209A-15

Query Match      88.0%; Score 673; DB 15; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQPWEHVNAIQEARRLLNLSRDTAAENNETVEVISEMFDLQEPCTCLOTRLE 75
Db 1 APARSPSPSTQPWEHVNAIQEARRLLNLSRDTAAENNETVEVISEMFDLQEPCTCLOTRLE 60

QY 76 LYKQGLRGSITKLGPLTWMAHYKQHCPTTPTSCATQIITPESPKENLKDFLLVIPP 135
Db 61 LYKQGLRGSITKLGPLTWMAHYKQHCPTTPTSCATQIITPESPKENLKDFLLVIPP 120

QY 136 CWPVQOE 142
Db 121 CWPVQOE 127
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```

RESULT 13
US-10-743-295-5
; Sequence 5, Application US/10743295
; Publication No. US20040136952A1
; GENERAL INFORMATION:
; APPLICANT: Shaekaran, Shyam S.
; APPLICANT: Sherman, Merry R.
; APPLICANT: Saifer, Mark G.P.
; APPLICANT: Williams, L. David
; TITLE OF INVENTION: POLYMER CONJUGATES OF CYTOKINES, CHEMOKINES, GROWTH FACTORS, POLY
; TITLE OF INVENTION: HORMONES AND ANTAGONISTS THEREOF WITH PRESERVED RECEPTOR-BINDING
; TITLE OF INVENTION: ACTIVITY
; FILE REFERENCE: 2057.0060002/JAG/BJD
; CURRENT FILING DATE: 2003-12-23
; PRIOR FILING DATE: 2003-12-23
; PRIOR FILING DATE: 2003-06-20
; PRIOR FILING DATE: 2002-12-26
; NUMBER OF SEQ ID NOS: 9
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5
; LENGTH: 127
; TYPE: PRT
; ORGANISM: Homo sapiens

Query Match      88.0%; Score 673; DB 16; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAENNETVEISEMFDLQEPCTCLOTRLE 75
DB 1 APARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAENNETVEISEMFDLQEPCTCLOTRLE 60

QY 76 LYKQGLRGSITKLGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPFD 135
DB 61 LYKQGLRGSITKLGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPFD 120

QY 136 CWEPVQE 142
DB 121 CWEPVQE 127

RESULT 14
US-10-743-295-5
; Sequence 5, Application US/10743295
; Publication No. US20040136952A1
; GENERAL INFORMATION:
; APPLICANT: Shaekaran, Shyam S.
; APPLICANT: Sherman, Merry R.
; APPLICANT: Saifer, Mark G.P.
; APPLICANT: Williams, L. David
; TITLE OF INVENTION: POLYMER CONJUGATES OF CYTOKINES, CHEMOKINES, GROWTH FACTORS, POLY
; TITLE OF INVENTION: HORMONES AND ANTAGONISTS THEREOF WITH PRESERVED RECEPTOR-BINDING
; TITLE OF INVENTION: ACTIVITY
; FILE REFERENCE: 2057.0060002/JAG/BJD
; CURRENT FILING DATE: 2003-12-23
; PRIOR FILING DATE: 2003-12-23
; PRIOR FILING DATE: 2003-06-20
; PRIOR FILING DATE: 2002-12-26
; NUMBER OF SEQ ID NOS: 9
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 5
; LENGTH: 127
; TYPE: PRT
; ORGANISM: Homo sapiens

Query Match      88.0%; Score 673; DB 16; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAENNETVEISEMFDLQEPCTCLOTRLE 75
DB 1 APARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAENNETVEISEMFDLQEPCTCLOTRLE 60

QY 76 LYKQGLRGSITKLGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPFD 135
DB 61 LYKQGLRGSITKLGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPFD 120

QY 136 CWEPVQE 142
DB 121 CWEPVQE 127

RESULT 15
US-10-773-939-8
; Sequence 8, Application US/10773939
; Publication No. US20040175356A1
; GENERAL INFORMATION:
; APPLICANT: Cox III, George N
; APPLICANT: Bolger Biotechnology, Inc.
; TITLE OF INVENTION: Derivatives of Growth Hormone and Related Proteins
; FILE REFERENCE: 4152-1-PUS
; CURRENT FILING DATE: 2004-02-05
; PRIOR FILING DATE: 2003-03-26
; PRIOR FILING DATE: 2003-03-26
; PRIOR FILING DATE: 2000-01-14
; PRIOR FILING DATE: 1997-07-14
; NUMBER OF SEQ ID NOS: 41
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 8
; LENGTH: 127
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-773-939-8

Query Match      88.0%; Score 673; DB 16; Length 127;
Best Local Similarity 100.0%; Pred. No. 2.5e-63;
Matches 127; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 16 APARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAENNETVEISEMFDLQEPCTCLOTRLE 75
DB 1 APARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAENNETVEISEMFDLQEPCTCLOTRLE 60

QY 76 LYKQGLRGSITKLGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPFD 135
DB 61 LYKQGLRGSITKLGPLTMASHYKQHCPTPTSCATQIITFESFKENLKDFLLVIPFD 120

QY 136 CWEPVQE 142
DB 121 CWEPVQE 127

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Search completed: March 8, 2005, 16:26:00  
Job time : 132 secs

Result No.	Score	Query Match	Length	DB	ID	Description	
1	682	89.2	259	3	US-08-469-318-141	Sequence 141, App	
2	682	89.2	259	3	US-08-468-609A-141	Sequence 141, App	
3	682	89.2	259	3	US-08-446-872A-141	Sequence 141, App	
4	682	89.2	259	4	US-08-762-227A-141	Sequence 141, App	
5	682	89.2	259	5	PCT-US95-01185-141	Sequence 141, App	
6	678	88.6	128	3	US-08-469-318-160	Sequence 160, App	
7	678	88.6	128	3	US-08-468-609A-160	Sequence 160, App	
8	678	88.6	128	3	US-08-446-872A-160	Sequence 160, App	
9	678	88.6	128	4	US-08-762-227A-160	Sequence 160, App	
10	678	88.6	128	5	PCT-US95-01185-160	Sequence 160, App	
11	678	88.6	274	3	US-08-469-318-144	Sequence 144, App	
12	678	88.6	274	3	US-08-468-609A-144	Sequence 144, App	
13	678	88.6	274	3	US-08-446-872A-144	Sequence 144, App	
14	678	88.6	274	4	US-08-762-227A-144	Sequence 144, App	
15	678	88.6	274	5	PCT-US95-01185-144	Sequence 144, App	
16	678	88.6	301	3	US-08-469-318-142	Sequence 142, App	
17	678	88.6	301	3	US-08-468-609A-142	Sequence 142, App	
18	678	88.6	301	3	US-08-446-872A-142	Sequence 142, App	
19	678	88.6	301	4	US-08-762-227A-142	Sequence 142, App	
20	678	88.6	301	5	PCT-US95-01185-142	Sequence 142, App	
21	673	88.0	127	1	US-08-318-193-2	Sequence 2, Appli	
22	673	88.0	127	4	US-09-462-941-8	Sequence 8, Appli	
23	673	88.0	127	6	5229496-15	Patent No. 5229496	
24	673	88.0	127	6	5229496-15	Patent No. 5229496	
25	673	88.0	131	6	5229496-2	Patent No. 5229496	
26	673	88.0	131	6	5229496-2	Patent No. 5229496	
27	673	88.0	144	1	US-08-284-393B-11	Sequence 11, Appl	

Patent No. 6030812  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; APPLICANT: Bauer, S. C.  
; APPLICANT: Braford-Goldberg, Sarah R.  
; APPLICANT: Caparon, Mair H.  
; APPLICANT: Easton, Alan M.  
; APPLICANT: Klein, Barbara K.  
; APPLICANT: McKeane, John P.  
; APPLICANT: Olin, Peter O.  
; APPLICANT: Paik, Kuman  
; APPLICANT: Thomas, John W.  
; TITLE OF INVENTION: Fusion Proteins Comprising Multiply Mutated Inteleukin-3 (III-3)  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
; ADDRESSEE: Corporate Patent Dept.  
; STREET: P. O. Box 5110  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60680  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/468,609A  
; FILING DATE: 06-JUN-1995  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Bennett, Dennis A.  
; REGISTRATION NUMBER: 34,547  
; REFERENCE/DOCKET NUMBER: C-2790/3  
; TELEPHONE: (314) 737-6986  
; TELEFAX: (314) 737-6972  
; INFORMATION FOR SEQ ID NO: 141:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 259 amino acids  
; TYPE: amino acid  
; STRANDEDNESS:  
; TOPOLOGY: linear  
; MOLECULE TYPE: protein  
; US-08-468-609A-141

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Best Local Similarity 97.0%; Pred. No. 1e-69;  
Matches 130; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
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Db 126 SGGGNNAPARSPSTQPEWHVNAIQEARRLLNLSRDTAENNETVEVISEMFDIQEPT 185  
QY 69 CLOTRLELYKQGLRSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLQDF 128  
Db 186 CLOTRLELYKQGLRSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLQDF 245  
QY 129 LLVIPDCWEPVQE 142  
Db 246 LLVIPDCWEPVQE 259

RESULT 3  
US-08-446-872A-141  
; Sequence 141, Application US/08446872A  
; Patent No. 6361977  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.

APPLICANT: Bauer, S. C.  
; APPLICANT: Braford-Goldberg, Sarah R.  
; APPLICANT: Caparon, Mair H.  
; APPLICANT: Easton, Alan M.  
; APPLICANT: Klein, Barbara K.  
; APPLICANT: McKeane, John P.  
; APPLICANT: Olin, Peter O.  
; APPLICANT: Paik, Kuman  
; APPLICANT: Thomas, John W.  
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis  
; TITLE OF INVENTION: Fusion Protein  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
; ADDRESSEE: Corporate Patent Dept.  
; STREET: P. O. Box 5110  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60680  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/446,872A  
; FILING DATE: 06-JUN-1995  
; CLASSIFICATION: 424  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Bennett, Dennis A.  
; REGISTRATION NUMBER: 34,547  
; REFERENCE/DOCKET NUMBER: C-2790/1  
; TELEPHONE: (314) 737-6986  
; TELEFAX: (314) 737-6972  
; INFORMATION FOR SEQ ID NO: 141:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 259 amino acids  
; TYPE: amino acid  
; STRANDEDNESS:  
; TOPOLOGY: linear  
; MOLECULE TYPE: protein  
; US-08-446-872A-141

Query Match 89.2%; Score 682; DB 3; Length 259;  
Best Local Similarity 97.0%; Pred. No. 1e-69;  
Matches 130; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 9 SGIEGMAPARSPSTQPEWHVNAIQEARRLLNLSRDTAENNETVEVISEMFDIQEPT 68  
Db 126 SGGGNNAPARSPSTQPEWHVNAIQEARRLLNLSRDTAENNETVEVISEMFDIQEPT 185  
QY 69 CLOTRLELYKQGLRSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLQDF 128  
Db 186 CLOTRLELYKQGLRSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLQDF 245  
QY 129 LLVIPDCWEPVQE 142  
Db 246 LLVIPDCWEPVQE 259

RESULT 4  
US-08-762-227A-141  
; Sequence 141, Application US/08762227A  
; Patent No. 6436387  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; Bauer, S. C.  
; Braford-Goldberg, Sarah R.

```
; Caparon, Mairé H.
; Easton, Alan M.
; Klein, Barbara K.
; McKearn, John P.
; Olin, Peter O.
; Paik, Kumnan
; Thomas, John W.
;
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis
; Fusion Protein
;
; NUMBER OF SEQUENCES: 197
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,
; STREET: P. O. Box 5110
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60680
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/762,227A
; FILING DATE: 09-Dec-1996
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/192,325
; FILING DATE: 14-FEB-1994
; APPLICATION NUMBER: US 08/446,872
; FILING DATE: 06-JUN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Bennett, Dennis A.
; REGISTRATION NUMBER: 34,547
; REFERENCE/DOCKET NUMBER: C-2790/5
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (708)470-6501
; TELEFAX: (708)470-6881
; INFORMATION FOR SEQ ID NO: 141:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 259 amino acids
; TYPE: amino acid
; STRANDEDNESS: <Unknown>
; TOPOLOGY: linear
; MOLECULE TYPE: protein
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; US-08-762-227A-141
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; Query Match 89.2%; Score 682; DB 4; Length 259;
; Best Local Similarity 97.0%; Pred. No. 1e-69;
; Matches 130; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
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; DB 126 SGGGNNAPSPSPSTQPEWHVNAIQEARRLLNLSRDTAANNETVEVISEMFDLQEPT 185
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; QY 69 CLQTRLELYKQGLRSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLKDF 128
; DB 186 CLQTRLELYKQGLRSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLKDF 245
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; QY 129 LLVIPDCWEPVQE 142
; DB 246 LLVIPDCWEPVQE 259
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; RESULT 5
; US-08-762-227A-141
; Sequence 141, Application PC/TUS9501185
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion
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; NUMBER OF SEQUENCES: 197
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/469,318
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/446,872
; FILING DATE:
; INFORMATION FOR SEQ ID NO: 160:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 128 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-469-318-160
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; Query Match 88.6%; Score 678; DB 3; Length 128;
; Best Local Similarity 100.0%; Pred. No. 1.1e-69;
; Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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; QY 9 SGIEGRMAPSPSPSTQPEWHVNAIQEARRLLNLSRDTAANNETVEVISEMFDLQEPT 68
; DB 126 SGGGNNAPSPSPSTQPEWHVNAIQEARRLLNLSRDTAANNETVEVISEMFDLQEPT 185
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; QY 69 CLQTRLELYKQGLRSLTKLKGPLTWASHYKQHCPTPTSCATQIITFESFKNLKDF 128
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; QY 129 LLVIPDCWEPVQE 142
; DB 246 LLVIPDCWEPVQE 259
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; RESULT 6
; US-08-469-318-160
; Sequence 160, Application US/08469318
; Patent No. 6022535
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion
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; NUMBER OF SEQUENCES: 196
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/469,318
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/446,872
; FILING DATE:
; INFORMATION FOR SEQ ID NO: 160:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 128 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-469-318-160
;
; Query Match 88.6%; Score 678; DB 3; Length 128;
; Best Local Similarity 100.0%; Pred. No. 1.1e-69;
; Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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61	ELYKGLRGSLTKLGPPLTMASHYKHCPTPTETSCATQIIIFESFKNLKDFFLVIIPF	120
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RESULT 7
US-08-468-609A-160
; Sequence 160, Application US/08468609A
; Patent No. 6030812
; GENERAL INFORMATION:
; APPLICANT: Abrams, Mark A.
; APPLICANT: Bauer, S. C.
; APPLICANT: Braford-Goldberg, Sarah R.
; APPLICANT: Caparon, Mairé H.
; APPLICANT: Easton, Alan M.
; APPLICANT: Klein, Barbara K.
; APPLICANT: McKearn, John P.
; APPLICANT: Olin, Peter O.
; APPLICANT: Paik, Kumhan
; APPLICANT: Thomas, John W.
; TITLE OF INVENTION: Fusion Proteins Comprising Multiply Mutated Inteleukin-3 (IL-
; NUMBER OF SEQUENCES: 197
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,
; ADDRESSEE: Corporate Patent Dept.
; STREET: P. O. Box 5110
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60680
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/468,609A
; FILING DATE: 06-JUN-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/192,325
; FILING DATE: 14-FEB-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Bennett, Dennis A.
; REGISTRATION NUMBER: 34,547
; REFERENCE/DOCKET NUMBER: C-2790/3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)737-6986
; TELEFAX: (314)737-6972
; INFORMATION FOR SEQ ID NO: 160:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 128 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-468-609A-160

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Query Match      88.6%; Score 678; DB 3; Length 128;
Best Local Similarity 100.0%; Pred. NO. 1.1e-69;
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0

QY 15 MAPARSPSPQTPWEHVNAIQEARRLLNLRDRTAAEENETVEISSMFDLOEPTCLQTRL 74
Db 1 MAPARSPSPQTPWEHVNAIQEARRLLNLRDRTAAEENETVEISSMFDLOEPTCLQTRL 60
OY 75 ELYKQGLRGSLTKLKGPLTMMASHYKHQCHPPTPETS CATOIIFESFKENLKDFLLIVIFP 134

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Db 61 ELYKQGLRSLTKLKGPLTMMASHYKQHCPTTSCATQIITFESFKNLKDPLLVIPT 120  
QY 135 DCWEPVQE 142  
Db 121 DCWEPVQE 128

## RESULT 9

US-08-762-227A-160  
; Sequence 160, Application US/08762227A  
; Patent No. 6436387  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; Bauer, S. C.  
; Bradford-Goldberg, Sarah R.  
; Caparon, Maïre H.  
; Easton, Alan M.  
; Klein, Barbara K.  
; McKearn, John P.  
; Olin, Peter O.  
; Paik, Kumnan  
; Thomas, John W.  
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis  
; FUSION PROTEIN  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
; Corporate Patent Dept.  
; STREET: P. O. Box 5110  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60680

COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/762,227A  
; FILING DATE: 09-Dec-1996

CLASSIFICATION: <Unknown>  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192,325

FILING DATE: 14-FEB-1994  
; APPLICATION NUMBER: US 08/446,872

FILING DATE: 06-JUN-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Bennett, Dennis A.

REGISTRATION NUMBER: 34,547  
; REFERENCE/DOCKET NUMBER: C-2790/5

TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (708) 470-6501

TELEFAX: (708) 470-6881  
; INFORMATION FOR SEQ ID NO: 160:

SEQUENCE CHARACTERISTICS:  
; LENGTH: 128 amino acids

TYPE: amino acid  
; STRANDEDNESS: <Unknown>

TOPOLOGY: linear  
; MOLECULE TYPE: protein

SEQUENCE DESCRIPTION: SEQ ID NO: 160:  
US-08-762-227A-160

Query Match 88.6%; Score 678; DB 4; Length 128;  
; Best Local Similarity 100.0%; Pred. No. 1.1e-69;  
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPQWEHVNAIQEARRLLNLSRDTAAEMNETVEVISEMFDLQPTCLOTRL 74  
Db 1 MAPARSPSPQWEHVNAIQEARRLLNLSRDTAAEMNETVEVISEMFDLQPTCLOTRL 60

QY 75 ELYKQGLRSLTKLKGPLTMMASHYKQHCPTTSCATQIITFESFKNLKDPLLVIPT 134  
Db 61 ELYKQGLRSLTKLKGPLTMMASHYKQHCPTTSCATQIITFESFKNLKDPLLVIPT 120  
QY 135 DCWEPVQE 142  
Db 121 DCWEPVQE 128

## RESULT 10

PCT-US95-01185-160  
; Sequence 160, Application PC/TUS9501185  
; GENERAL INFORMATION:

; APPLICANT:

; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion

; TITLE OF INVENTION: Protein

; NUMBER OF SEQUENCES: 196

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent In Release #1.0, Version #1.30 (EPO)

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: PCT/US95/01185

; FILING DATE: 02-FEB-1995

; CLASSIFICATION:

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 08/192325

; FILING DATE: 14-FEB-1994

; INFORMATION FOR SEQ ID NO: 160:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 128 amino acids

; TYPE: amino acid

; STRANDEDNESS:

; TOPOLOGY: linear

; MOLECULE TYPE: protein

PCT-US95-01185-160

Query Match 88.6%; Score 678; DB 5; Length 128;

Best Local Similarity 100.0%; Pred. No. 1.1e-69;

Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPQWEHVNAIQEARRLLNLSRDTAAEMNETVEVISEMFDLQPTCLOTRL 74

Db 1 MAPARSPSPQWEHVNAIQEARRLLNLSRDTAAEMNETVEVISEMFDLQPTCLOTRL 60

QY 75 ELYKQGLRSLTKLKGPLTMMASHYKQHCPTTSCATQIITFESFKNLKDPLLVIPT 134

Db 61 ELYKQGLRSLTKLKGPLTMMASHYKQHCPTTSCATQIITFESFKNLKDPLLVIPT 120

QY 135 DCWEPVQE 142

Db 121 DCWEPVQE 128

## RESULT 11

US-08-469-318-144

; Sequence 144, Application US/08469318

; Patent No. 6022535

; GENERAL INFORMATION:

; APPLICANT:

; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion

; TITLE OF INVENTION: Protein

; NUMBER OF SEQUENCES: 196

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: Patent In Release #1.0, Version #1.30 (EPO)

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/469,318

; FILING DATE:

; CLASSIFICATION:

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/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: 08/446,872
/ FILING DATE:
/ INFORMATION FOR SEQ ID NO: 144:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 274 amino acids
/ TYPE: amino acid
/ STRANDEDNESS:
/ TOPOLOGY: linear
/ MOLECULE TYPE: protein
/ US-08-469-318-144

Query Match      88.6%; Score 678; DB 3; Length 274;
Best Local Similarity 100.0%; Pred. No. 3.2e-69;
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAEMNETVEVISMFQLOEPTCLOTRL 74
DB 147 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAEMNETVEVISMFQLOEPTCLOTRL 206
QY 75 ELYKQGLRGLTKLKGPLTMASHYKQHCPPPTPETS CATQIITFESFKENLKDFLVIPP 134
DB 207 ELYKQGLRGLTKLKGPLTMASHYKQHCPPPTPETS CATQIITFESFKENLKDFLVIPP 266
QY 135 DCWEPVQE 142
DB 267 DCWEPVQE 274

RESULT 12
US-08-469-609A-144
/ Sequence 144, Application US/08468609A
/ Patent No. 6030812
/ GENERAL INFORMATION:
/ APPLICANT: Abrams, Mark A.
/ APPLICANT: Bauer, S. C.
/ APPLICANT: Braford-Goldberg, Sarah R.
/ APPLICANT: Caparon, Maïre H.
/ APPLICANT: Easton, Alan M.
/ APPLICANT: Klein, Barbara K.
/ APPLICANT: McKearn, John P.
/ APPLICANT: Olin, Peter O.
/ APPLICANT: Paik, Kuman
/ APPLICANT: Thomas, John W.
/ APPLICANT: Thomas, John W.
/ TITLE OF INVENTION: Fusion Proteins Comprising Multiply Mutated Inteleukin-3 (IL-
/ NUMBER OF SEQUENCES: 197
/ CORRESPONDENCE ADDRESS:
/ ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,
/ ADDRESSEE: Corporate Patent Dept.
/ STREET: P. O. Box 5110
/ CITY: Chicago
/ STATE: Illinois
/ COUNTRY: USA
/ ZIP: 60680
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Floppy disk
/ COMPUTER: IBM PC compatible
/ OPERATING SYSTEM: PC-DOS/MS-DOS
/ SOFTWARE: Patent In Release #1.0, Version #1.25
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/08/469,609A
/ FILING DATE: 06-JUN-1995
/ CLASSIFICATION:
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: US 08/192,325
/ FILING DATE: 14-FEB-1994
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Bennett, Dennis A.
/ REGISTRATION NUMBER: 34,547
/ REFERENCE/DOCKET NUMBER: C-2790/3
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: (314)737-6986
/ TELEFAX: (314)737-6972
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/ INFORMATION FOR SEQ ID NO: 144:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 274 amino acids
/ TYPE: amino acid
/ STRANDEDNESS:
/ TOPOLOGY: linear
/ MOLECULE TYPE: protein
/ US-08-468-609A-144

Query Match      88.6%; Score 678; DB 3; Length 274;
Best Local Similarity 100.0%; Pred. No. 3.2e-69;
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAEMNETVEVISMFQLOEPTCLOTRL 74
DB 147 MAPARSPSPSTQPEWHVNAIQEARRLLNLSRDTAAEMNETVEVISMFQLOEPTCLOTRL 206
QY 75 ELYKQGLRGLTKLKGPLTMASHYKQHCPPPTPETS CATQIITFESFKENLKDFLVIPP 134
DB 207 ELYKQGLRGLTKLKGPLTMASHYKQHCPPPTPETS CATQIITFESFKENLKDFLVIPP 266
QY 135 DCWEPVQE 142
DB 267 DCWEPVQE 274

RESULT 13
US-08-446-872A-144
/ Sequence 144, Application US/08446872A
/ Patent No. 6361977
/ GENERAL INFORMATION:
/ APPLICANT: Abrams, Mark A.
/ APPLICANT: Bauer, S. C.
/ APPLICANT: Braford-Goldberg, Sarah R.
/ APPLICANT: Caparon, Maïre H.
/ APPLICANT: Easton, Alan M.
/ APPLICANT: Klein, Barbara K.
/ APPLICANT: McKearn, John P.
/ APPLICANT: Olin, Peter O.
/ APPLICANT: Paik, Kuman
/ APPLICANT: Thomas, John W.
/ APPLICANT: Thomas, John W.
/ TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis
/ NUMBER OF SEQUENCES: 197
/ CORRESPONDENCE ADDRESS:
/ ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,
/ ADDRESSEE: Corporate Patent Dept.
/ STREET: P. O. Box 5110
/ CITY: Chicago
/ STATE: Illinois
/ COUNTRY: USA
/ ZIP: 60680
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Floppy disk
/ COMPUTER: IBM PC compatible
/ OPERATING SYSTEM: PC-DOS/MS-DOS
/ SOFTWARE: Patent In Release #1.0, Version #1.25
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/08/446,872A
/ FILING DATE: 06-JUN-1995
/ CLASSIFICATION:
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: US 08/192,325
/ FILING DATE: 14-FEB-1994
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Bennett, Dennis A.
/ REGISTRATION NUMBER: 34,547
/ REFERENCE/DOCKET NUMBER: C-2790/1
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: (314)737-6986
/ TELEFAX: (314)737-6972
/ INFORMATION FOR SEQ ID NO: 144:
/ SEQUENCE CHARACTERISTICS:
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;
; LENGTH: 274 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-446-872A-144
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; Query Match 88.6%; Score 678; DB 3; Length 274;
; Best Local Similarity 100.0%; Pred. No. 3.2e-69;
; Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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; QY 15 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAEMNETVEVISEMFDLQEPFCLQTRL 74
; DB 147 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAEMNETVEVISEMFDLQEPFCLQTRL 206
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; QY 75 ELYKQGLRGLSLTKLKGPLTMMASHYKQHCPPTPETSATQIITFESPKENLKDPLLVIPF 134
; DB 207 ELYKQGLRGLSLTKLKGPLTMMASHYKQHCPPTPETSATQIITFESPKENLKDPLLVIPF 266
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; QY 135 DCWEPVQE 142
; DB 267 DCWEPVQE 274
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; RESULT 14
; US-08-762-227A-144
; Sequence 144, Application US/08762227A
; Patent No. 6436387
; GENERAL INFORMATION:
; APPLICANT: Abrams, Mark A.
; Bratford-Goldberg, Sarah R.
; Caparon, Maire H.
; Easton, Alan M.
; Klein, Barbara K.
; Mckearn, John P.
; Oline, Peter O.
; Paik, Kumnan
; Thomas, John W.
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis
; FUSION PROTEIN
; NUMBER OF SEQUENCES: 197
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,
; Corporate Patent Dept.
; STREET: P. O. Box 5110
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60680
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/762,227A
; FILING DATE: 09-Dec-1996
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/192,325
; FILING DATE: 14-FEB-1994
; APPLICATION NUMBER: US 08/446,872
; FILING DATE: 06-JUN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Bennett, Dennis A.
; REGISTRATION NUMBER: 34,547
; REFERENCE/DOCKET NUMBER: C-2790/5
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (708)470-6501
; TELEFAX: (708)470-6881
; INFORMATION FOR SEQ ID NO: 144:
; SEQUENCE CHARACTERISTICS:
;
; LENGTH: 274 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-446-872A-144
;
; Query Match 88.6%; Score 678; DB 3; Length 274;
; Best Local Similarity 100.0%; Pred. No. 3.2e-69;
; Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
;
; QY 15 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAEMNETVEVISEMFDLQEPFCLQTRL 74
; DB 147 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAEMNETVEVISEMFDLQEPFCLQTRL 206
;
; QY 75 ELYKQGLRGLSLTKLKGPLTMMASHYKQHCPPTPETSATQIITFESPKENLKDPLLVIPF 134
; DB 207 ELYKQGLRGLSLTKLKGPLTMMASHYKQHCPPTPETSATQIITFESPKENLKDPLLVIPF 266
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; QY 135 DCWEPVQE 142
; DB 267 DCWEPVQE 274
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; RESULT 15
; PCT-US95-01185-144
; Sequence 144, Application PC/TUS9501185
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion
; TITLE OF INVENTION: Protein
; NUMBER OF SEQUENCES: 196
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30 (BPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/01185
; FILING DATE: 02-FEB-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/192325
; FILING DATE: 14-FEB-1994
; INFORMATION FOR SEQ ID NO: 144:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 274 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; PCT-US95-01185-144
;
; Query Match 88.6%; Score 678; DB 5; Length 274;
; Best Local Similarity 100.0%; Pred. No. 3.2e-69;
; Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
;
; QY 15 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAEMNETVEVISEMFDLQEPFCLQTRL 74
; DB 147 MAPARSPSTQPEWHVNAIQEARRLLNLSRDAAEMNETVEVISEMFDLQEPFCLQTRL 206
;
; QY 75 ELYKQGLRGLSLTKLKGPLTMMASHYKQHCPPTPETSATQIITFESPKENLKDPLLVIPF 134
; DB 207 ELYKQGLRGLSLTKLKGPLTMMASHYKQHCPPTPETSATQIITFESPKENLKDPLLVIPF 266
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; QY 135 DCWEPVQE 142
; DB 267 DCWEPVQE 274
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GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: March 11, 2005, 10:03:16 ; Search time 2574 Seconds  
(without alignments)  
8621.791 Million cell updates/sec

Title: US-10-723-083-1  
Perfect score: 458  
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Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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2: gb\_hgt.\*  
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6: gb\_pat.\*  
7: gb\_ph.\*  
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14: gb\_vi.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	319.2	69.7	909	6	I49839	I49839 Sequence 9
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17	283	61.8	402	6	AR223282	AR223282 Sequence
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19	281	61.4	822	6	AR223222	AR223222 Sequence

20	281	61.4	903	6	AR202217	AR202217 Sequence
21	281	61.4	903	6	AR223219	AR223219 Sequence
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C 23	277.6	60.6	415	6	A00368	A00368 Artificial
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C 25	277.6	60.6	415	6	A14306	A14306 GM-CSF gene
26	276.8	60.4	756	6	CQ721607	CQ721607 Sequence
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33	276.8	60.4	787	6	I09160	I09160 Sequence 21
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35	276.8	60.4	2385	6	AR082744	AR082744 Sequence
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38	276.2	60.3	644	6	E01141	E01141 cDNA encodi
39	276.2	60.3	660	6	AR364645	AR364645 Sequence
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41	276.2	60.3	1011	6	AR447690	AR447690 Sequence
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ALIGNMENTS

RESULT 1  
LOCUS A20088 905 bp DNA linear PAT 18-AUG-1994  
DEFINITION BamHI-HindIII fragment in vector pAEO.GMCSF.  
ACCESSION A20088  
VERSION A20088.1 GI:583268  
KEYWORDS synthetic construct  
SOURCE synthetic construct  
ORGANISM other sequences; artificial sequences.  
REFERENCE 1 (bases 1 to 905)  
AUTHORS Garvin,R.T. and Malek,L.T.  
TITLE An expression system for the secretion of bioactive human granulocyte macrophage colony stimulating factor (GM-CSF) and other heterologous proteins from streptomyces  
JOURNAL Patent: EP 0352707-A 21 31-JAN-1990;  
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ORIGIN

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Best Local Similarity 86.9%; Pred. No. 3.6e-38;  
Matches 351; Conservative 0; Mismatches 53; Indels 0; Gaps 0;  
Qy 35 CCGGATCGAGGGCGCATGGCGCCAGCGCGAGCCCGGCGGTCACCCAGCCGCTGGG 94  
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Qy 95 AGCAGCTGAACCGCATCCAGGAGGCGCGGAGGCTCTCAACTCTCTCCCGGACCGCGCG 154  
Db 562 AGCAGCTGAACCGCATCCAGGAGGCGCGGAGGCTCTCAACTCTCTCGGGGACCGCGCG 621  
Qy 155 CCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTCCGATCTCCAGGAGCCGACCT 214  
Db 622 CCGAGATGAACGAGACCGTGGAGGTGATCTCGGAGATGTTCCGACTTTCAGGAGCCACGT 681  
Qy 215 GCCTCAGACCCGCTCGAGCTGTACAGCAGGGGCTCGCGGGAGCCCTCACCAGCTCA 274

Db 682 GCCTCCAGACCGCCTCGAGCTGTATCAAGACAGGGGCTCCGGGCGACCTCCACCAAGCTCA 741

QY 275 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 334

Db 742 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 801

QY 335 CCTCTCGGCGCCACCCAGATCATCACTTCGAGAGCTTCAAGGAGAACTCAAGGACTTCC 394

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RESULT 2  
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LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
FEATURES  
source

AR363245  
Sequence 3 from patent US 5200327.  
AR363245  
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SOURCE  
ORGANISM  
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REFERENCE  
1 (bases 1 to 905)  
Garvin,R.T. and Malek,L.T.  
Expression system for the secretion of bioactive human granulocyte macrophage colony stimulating factor (GM-CSF) and other heterologous proteins from streptomyces  
Patent: US 5200327-A 3 06-APR-1993;  
JOURNAL  
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Best Local Similarity 86.9%; Pred. No. 3.6e-38;  
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QY 35 CCGGCATCGAGGCGCGATGGCCAGCGCGAGCCCGGCTCCACCCAGCGGTGGG 94

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QY 95 AGCAGTGAACGCGATCCAGAGCGCGGAGGCTCTCAACCTCTCCCGCAGACCGCGG 154

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QY 155 CCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTTCGATCTCCAGGAGCCGACCT 214

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QY 335 CCTCTCGGCGCCACCCAGATCATCACTTCGAGAGCTTCAAGGAGAACTCAAGGACTTCC 394

Db 802 COTGTGCCCGCCACCCAGATCATCACTTCGAGTCTTCAAGGAGAACTCAAGGACTTCC 861

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RESULT 3  
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LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
ORGANISM  
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TITLE  
JOURNAL  
FEATURES  
source

A20089  
Sequence 3 from patent US 5200327.  
A20089  
A20089.1 GI:34424298  
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SOURCE  
ORGANISM  
Unclassified.  
REFERENCE  
1 (bases 1 to 905)  
Garvin,R.T. and Malek,L.T.  
Expression system for the secretion of bioactive human granulocyte macrophage colony stimulating factor (GM-CSF) and other heterologous proteins from streptomyces  
Patent: US 5200327-A 3 06-APR-1993;  
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QY 95 AGCAGTGAACGCGATCCAGAGCGCGGAGGCTCTCAACCTCTCCCGCAGACCGCGG 154

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QY 155 CCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTTCGATCTCCAGGAGCCGACCT 214

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QY 215 GCCTCCAGACCCCGCTCGAGCTGTACAAGCAGGGCTCCCGGCGAGCTCCACCAAGCTCA 274

Db 682 GCCTCCAGACCCCGCTCGAGCTGTACAAGCAGGGCTCCCGGCGAGCTCCACCAAGCTCA 741

QY 275 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 334

Db 742 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 801

QY 335 CCTCTCGGCGCCACCCAGATCATCACTTCGAGAGCTTCAAGGAGAACTCAAGGACTTCC 394

Db 802 COTGTGCCCGCCACCCAGATCATCACTTCGAGTCTTCAAGGAGAACTCAAGGACTTCC 861

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Db 862 TCCTCGTGATCCCGTTCGACTCTCGGAGCCCGGTCCAGGAGTGA 905

RESULT 4  
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LOCUS  
DEFINITION  
ACCESSION  
VERSION  
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TITLE  
JOURNAL  
FEATURES  
source

A20089  
Sequence 9 from patent US 5641663.  
A20089  
A20089.1 GI:2472059  
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SOURCE  
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REFERENCE  
1 (bases 1 to 909)  
Garvin,R.T. and Malek,L.T.  
Expression system for the secretion of bioactive human granulocyte macrophage colony stimulating factor (GM-CSF) and other heterologous proteins from streptomyces  
Patent: US 5641663-A 9 24-JUN-1997;  
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QY 95 AGCAGTGAACGCGATCCAGAGCGCGGAGGCTCTCAACCTCTCCCGCAGACCGCGG 154

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QY 155 CCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTTCGATCTCCAGGAGCCGACCT 214

Db 288 CCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTTCGACTTTCGAGGAGCCACGT 229

QY 215 GCCTCCAGACCCCGCTCGAGCTGTACAAGCAGGGCTCCCGGCGAGCTCCACCAAGCTCA 274

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QY 275 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 334

Db 168 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 109

QY 335 CCTCTCGGCGCCACCCAGATCATCACTTCGAGAGCTTCAAGGAGAACTCAAGGACTTCC 394

Db 108 COTGTGCCCGCCACCCAGATCATCACTTCGAGTCTTCAAGGAGAACTCAAGGACTTCC 49

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DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
FEATURES  
source

BamHI-HindIII fragment in vector pABO.GMCSF.  
A20089  
A20089.1 GI:578985  
synthetic construct  
synthetic construct  
other sequences; artificial sequences.  
1 (bases 1 to 906)  
Garvin,R.T. and Malek,L.T.  
An expression system for the secretion of bioactive human granulocyte macrophage colony stimulating factor (GM-CSF) and other heterologous proteins from streptomyces  
Patent: EP 0352707-A 22 31-JAN-1990;  
Cangene Corporation  
Location/Qualifiers  
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QY 35 CCGGCATCGAGGCGCGATGGCCAGCGCGAGCCCGGCTCCACCCAGCGGTGGG 94

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QY 95 AGCAGTGAACGCGATCCAGAGCGCGGAGGCTCTCAACCTCTCCCGCAGACCGCGG 154

Db 348 AGCAGTGAACGCGATCCAGAGCGCGGAGGCTCTCAACCTCTCGCGGAGACCGCGG 289

QY 155 CCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTTCGATCTCCAGGAGCCGACCT 214

Db 288 CCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTTCGACTTTCGAGGAGCCACGT 229

QY 215 GCCTCCAGACCCCGCTCGAGCTGTACAAGCAGGGCTCCCGGCGAGCTCCACCAAGCTCA 274

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QY 275 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 334

Db 168 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 109

QY 335 CCTCTCGGCGCCACCCAGATCATCACTTCGAGAGCTTCAAGGAGAACTCAAGGACTTCC 394

Db 108 COTGTGCCCGCCACCCAGATCATCACTTCGAGTCTTCAAGGAGAACTCAAGGACTTCC 49

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RESULT 4  
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LOCUS  
DEFINITION  
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VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
FEATURES  
source

A20089  
Sequence 9 from patent US 5641663.  
A20089  
A20089.1 GI:2472059  
Unknown.  
SOURCE  
ORGANISM  
Unclassified.  
REFERENCE  
1 (bases 1 to 909)  
Garvin,R.T. and Malek,L.T.  
Expression system for the secretion of bioactive human granulocyte macrophage colony stimulating factor (GM-CSF) and other heterologous proteins from streptomyces  
Patent: US 5641663-A 9 24-JUN-1997;  
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FEATURES  
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Best Local Similarity 86.9%; Pred. No. 3.6e-38;  
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QY 35 CCGGCATCGAGGCGCGATGGCCAGCGCGAGCCCGGCTCCACCCAGCGGTGGG 94

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QY 95 AGCAGTGAACGCGATCCAGAGCGCGGAGGCTCTCAACCTCTCCCGCAGACCGCGG 154

Db 348 AGCAGTGAACGCGATCCAGAGCGCGGAGGCTCTCAACCTCTCGCGGAGACCGCGG 289

QY 155 CCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTTCGATCTCCAGGAGCCGACCT 214

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QY 275 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 334

Db 168 AGGGCCCGCTACCATGATGGGTCCCACTCAACAGCAGCAGTCCACCGACCCCGGAGA 109

QY 335 CCTCTCGGCGCCACCCAGATCATCACTTCGAGAGCTTCAAGGAGAACTCAAGGACTTCC 394

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QY 155 CCAGATGAACGAGACCGTGAGGTGATCTCCAGATGTTTCGATCTCCAGGAGCGACCT 214
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QY 215 GCCTCAGACCCGCTCGAGCTGTACAGAGGCGCTCCGGGAGCGCTTCAACGAGCTCA 274
DB 682 GCCTCAGACCCGCTCGAGCTGTACAGAGGCGCTCCGGGAGCGCTTCAACGAGCTCA 741
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RESULT 5
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LOCUS      384 bp DNA linear PAT 18-AUG-1994
DEFINITION PetI-Hind III fragment encoding GM-CSF.
ACCESSION A20082
VERSION A20082.1 GI:583266
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 384)
AUTHORS    Garvin,R.T. and Malek,L.T.
TITLE      An expression system for the secretion of bioactive human
            granulocyte macrophage colony stimulating factor (GM-CSF) and other
            heterologous proteins from streptomyces
            Patent: EP 0352707-A 13 31-JAN-1990;
            Cangene Corporation
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QY 115 GAGGCGCGAGGCTCTCAACCTCTCCCGGACACCGCGGAGATGAACGAGACCGTG 174
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RESULT 6
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LOCUS      392 bp DNA linear PAT 18-AUG-1994
DEFINITION PetI-Hind III fragment encoding GM-CSF.
ACCESSION A20083
VERSION A20083.1 GI:578981
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 392)
AUTHORS    Garvin,R.T. and Malek,L.T.
TITLE      An expression system for the secretion of bioactive human
            granulocyte macrophage colony stimulating factor (GM-CSF) and other
            heterologous proteins from streptomyces
            Patent: EP 0352707-A 14 31-JAN-1990;
            Cangene Corporation
FEATURES    Location/Qualifiers
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Best Local Similarity 89.3%; Pred. No. 5.6e-38;
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QY 55 GCGCCAGCGCGAGCCCGAGCCCGTCCACCCAGCGCGTGGAGACAGTGAACGCGATCCAG 114
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QY 115 GAGGCGCGAGGCTCTCAACCTCTCCCGGACACCGCGGAGATGAACGAGACCGTG 174
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Db 388 GCCCCGCGCGTCCGCTCGACCCAGCGTGGGAGCAGCTCAACGCGATCCAG 329  
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DEFINITION Sequence 2 from patent US 5200327.  
ACCESSION AR363244  
VERSION AR363244.1 GI:34424297  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE 1 (bases 1 to 896)  
AUTHORS Garvin,R.T. and Malek,L.T.  
TITLE Expression system for the secretion of bioactive human granulocyte macrophage colony stimulating factor (GM-CSF) and other heterologous proteins from streptomycetes  
JOURNAL Patent: US 5200327-A 2 06-APR-1993;  
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ORIGIN  
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Best Local Similarity 89.3%; Pred. No. 4.7e-38;  
Matches 343; Conservative 0; Mismatches 41; Indels 0; Gaps 0;  
QY 55 GCGCCAGCGGAGCGGCGGCTCCAGCCAGCGGAGGAGCAGCTGAACGCGATCCAG 114  
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LOCUS I49838 900 bp DNA linear PAT 07-OCT-1997  
DEFINITION Sequence 7 from patent US 5641663.  
ACCESSION I49838  
VERSION I49838.1 GI:2472058  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE 1 (bases 1 to 900)  
AUTHORS Garvin,R.T. and Malek,L.T.  
TITLE Expression system for the secretion of bioactive human granulocyte macrophage colony stimulating factor (GM-CSF) and other heterologous proteins from streptomycetes  
JOURNAL Patent: US 5641663-A 7 24-JUN-1997;  
FEATURES  
source Location/Qualifiers  
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Best Local Similarity 89.3%; Pred. No. 4.7e-38;  
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QY 115 GAGCCGCGAGGCTCTCTCAACCTCTCCCGGACACCGCCCGCAGATGAACGAGACCGTG 174  
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QY 175 GAGGTGATCTCCGAGATGTTTCGATCTCCAGAGCGGACCTGCTCCAGACCCGCTCGAG 234  
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QY 235 CTGTACAAGCAGGCGCTCCGCGGAGCCTCACAAGCTCAAGGCGCGCTCACCATGATG 294  
Db 693 CTGTACAAGCAGGCGCTCCGCGGAGCCTCACAAGCTCAAGGCGCGCTCACCATGATG 752  
QY 295 GCGTCCACTACAAGCAGCCTCCGCGGACACCGCCGAGATGAACGAGACCGTG 354  
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ACCESSION AR363243  
VERSION AR363243.1 GI:34424296  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.

Unclassified.									
REFERENCE	1 (bases 1 to 386)								
AUTHORS	Garvin,R.T. and Malek,L.T.								
TITLE	Expression system for the secretion of bioactive human granulocyte macrophage colony stimulating factor (GM-CSF) and other heterologous proteins from streptomyces								
JOURNAL	Patent: US 5200327-A 1 06-APR-1993;								
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QY	316 TGCCACCGACCCCGGAGACCTCTCTCGGCCACCCAGATCATCACTTCGAGAGCTTCAAG	375							
DB	264 TGCCCCCAGCGCGGAGAGCTGCTGGCCGCCACCATCATCACTTCGAGTGTTCAG	323							
QY	376 GAGAACTCAAGGACTTCTCTCTCGTGATCCCGTTCGACTGTGGGAGCGGTGCAGGAG	435							
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ACCESSION	CQ834915								
VERSION	CQ834915.1	GI:50834452							
KEYWORDS									
SOURCE	Homo sapiens (human)								
ORGANISM	Homo sapiens								
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.								
AUTHORS	1								
TITLE	Raab,D., Graf,M., Norka,F. and Wagner,R.								
JOURNAL	Method and device for optimizing a nucleotide sequence for the purpose of expression of a protein								
FEATURES	Patent: WO 2004059556-A 2 15-JUL-2004;								
source	Geneart GmbH (DE)								
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DEFINITION	Sequence 55 from patent US 6361977.								
ACCESSION	AR202206								
VERSION	AR202206.1	GI:20256745							
KEYWORDS									
SOURCE	Unknown.								
ORGANISM	Unknown.								
REFERENCE	1 (bases 1 to 777)								
AUTHORS	Bauer, S. Christopher., Abrams, M. Allen., Braford-Goldberg, S. Ruth., Caparon, M. Helena., Easton, A. Michael., Klein, B. Kure., McKearn, J. P., Olin, P. O., Paik, K. and Thomas, J. W.								
TITLE	Methods of using multivalent IL-3 hematopoiesis fusion protein								
JOURNAL	Patent: US 6361977-A 55 26-MAR-2002;								
FEATURES	Location/Qualifiers								
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QY	154 GCCGAGATGAACGAGACCGTGGAGTGATCTCCGAGATGTTGATCTCCAGGCGCGACC	213							
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Db 616 AAGGGCCCTTGACCATGATGGCCAGCCACTACAGCAGCACTGCCCTCCAAACCCCGGAA 675  
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## RESULT 15

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DEFINITION Sequence 55 from patent US 6436387.  
ACCESSION AR223208  
VERSION AR223208.1 GI:233331216

KEYWORDS  
SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 777)

AUTHORS Bauer, S.C., Abrams, M.A., Braford-Goldberg, S.P., Caparon, M.H.,  
Easton, A.M., Klein, B.K., McKearn, J.P., Olin, P.O., Paik, K. and  
Thomas, J.W.

TITLE Methods of ex-vivo expansion of hematopoietic cells using

JOURNAL multivariant IL-3 hematopoiesis chimera proteins

FEATURES Patent: US 6436387-A 55 20-AUG-2002;

source Location/Qualifiers

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/mol\_type="genomic DNA"

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Query Match 61.9%; Score 283.6; DB 6; Length 777;  
Best Local Similarity 81.8%; Pred. No. 7.1e-33;  
Matches 328; Conservative 0; Mismatches 74; Indels 0; Gaps 0;  
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Qy 154 GCCGAGATGAACAGACCGGTGGAGGTGATCTCCGAGATGTTGATCTCCAGAGCCGACC 213  
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Qy 214 TGCCTCCAGACCGCCTCGAGCTGTACAAGCAGGCGCTCCGCGCAGCCTCACCAAGCTC 273  
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GenCore version 5.1.6  
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

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Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

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Post-processing: Minimum Match 0%

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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

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2	319.2	69.7	905	6	5200327-3
3	319.2	69.7	909	1	US-08-318-193-9
4	318.4	69.5	392	1	US-08-318-193-1
5	318.4	69.5	896	6	5200327-2
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13	283.6	61.9	777	3	US-08-468-609A-55
14	283.6	61.9	777	3	US-08-446-872A-55
15	283.6	61.9	777	3	US-08-762-227A-55
16	283.6	61.9	777	3	PCT-US95-01185-55
17	283	61.8	402	3	US-08-469-318-176
18	283	61.8	402	3	US-08-468-609A-176
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31	281	61.4	903	5	PCT-US95-01185-66	Sequence 3, Appl
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34	276.8	60.4	2385	3	US-09-344-195-3	Sequence 3, Appl
35	276.2	60.3	660	6	5391485-2	Patent No. 5391485
36	276.2	60.3	660	6	5391485-2	Patent No. 5391485
37	276.2	60.3	661	6	5229496-1	Patent No. 5229496
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39	276.2	60.3	1011	4	US-09-976-594-275	Sequence 275, App
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43	275.2	60.1	1318	3	US-09-310-842-3	Sequence 3, Appl
44	274.6	60.0	1011	2	US-08-750-128-12	Sequence 12, Appl
45	274.6	60.0	1588	2	US-09-146-283-1	Sequence 1, Appl

#### ALIGNMENTS

RESULT 1

5200327-3

; Patent No. 5200327

; APPLICANT: GARVIN, ROBERT T.; MALEK, LAWRENCE T.

; TITLE OF INVENTION: EXPRESSION SYSTEM FOR THE SECRETION OF

; BIOACTIVE HUMAN GRANULOCYTE MACROPHAGE COLONY STIMULATING

; FACTOR (GM-CSF) AND OTHER HETEROLOGOUS PROTEINS FROM

; STREPTOMYCES

; NUMBER OF SEQUENCES: 24

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/07/224,568

; FILING DATE: 26-JUL-1988

; SEQ ID NO:3:

; LENGTH: 905

5200327-3

Query Match 69.7%; Score 319.2; DB 6; Length 905;  
Best Local Similarity 86.9%; Pred. No. 1e-60;  
Matches 351; Conservative: 0; Mismatches 53; Indels 0; Gaps 0;

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RESULT 2

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5200327-3
; Patent No. 5200327
; APPLICANT: GARVIN, ROBERT T.; MALEK, LAWRENCE T.
; TITLE OF INVENTION: EXPRESSION SYSTEM FOR THE SECRETION OF
; BIOACTIVE HUMAN GRANULOCYTE MACROPHAGE COLONY STIMULATING
; FACTOR (GM-CSF) AND OTHER HETEROLOGOUS PROTEINS FROM
; STREPTOMYCES
; NUMBER OF SEQUENCES: 24
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/224,568
; FILING DATE: 26-JUL-1988
; SEQ ID NO.: 3
; LENGTH: 905
5200327-3

Query Match 69.7%; Score 319.2; DB 6; Length 905;
Best Local Similarity 86.9%; Pred. No. 1e-60;
Matches 351; Conservative 0; Mismatches 53; Indels 0; Gaps 0;

Qy 35 CCGGCATCGAGGGCCGATGGCCAGCGGCGAGCCCGAGAGCCCGTCCACCCAGCGTGGG 94
Db 502 CCGCCTCCGGGGCGTCTGCAGCCCGCCCGGTCGCCGTCGCCGTCAGCCAGCGTGGG 561
Qy 95 AGCACGTGAACGGCATCCAGAGAGCCCGCAGGCTCCTCAACCTTCGCCGGACACCCGCG 154
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2  US-08-318-193-9
3  ; Sequence 9, Application US/08318193
4  ; Patent No. 5641663
5  ; GENERAL INFORMATION:
6  ; APPLICANT: GARVIN, Robert T.
7  ; APPLICANT: MALEK, Lawrence T.
8  ; TITLE OF INVENTION: AN EXPRESSION SYSTEM FOR THE SECRETION
9  ; TITLE OF INVENTION: OF BIOACTIVE HUMAN GRANULOCYTE MACROPHAGE COLONY
10 ; TITLE OF INVENTION: STIMULATING FACTOR (GM-CSF) AND OTHER HETEROLOGOUS
11 ; TITLE OF INVENTION: PROTEINS FROM STREPTOMYCES
12 ; NUMBER OF SEQUENCES: 91
13 ; CORRESPONDENCE ADDRESS:
14 ; ADDRESSEE: Foley & Lardner
15 ; STREET: 1900 Diagonal Road, Suite 500
16 ; CITY: Alexandria
17 ; STATE: Virginia
18 ; COUNTRY: USA
19 ; ZIP: 22313-0299
20 ; COMPUTER READABLE FORM:
21 ; MEDIUM TYPE: Floppy disk
22 ; COMPUTER: IBM PC compatible
23 ; OPERATING SYSTEM: PC-DOS/MS-DOS
24 ; SOFTWARE: PatentIn Release #1.0, Version #1.25
25 ; CURRENT APPLICATION DATA:

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; APPLICATION NUMBER: US/08/318,193
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/07/935,314
; FILING DATE:
; APPLICATION NUMBER: US 07/224,568
; ATTORNEY/AGENT INFORMATION:
; NAME: BENT, Stephen A.
; REGISTRATION NUMBER: 29,768
; REFERENCE/DOCKET NUMBER: 18740/116 CACO
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703)836-9300
; TELEFAX: (703)683-4109
; TELEX: 899149
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 909 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: Other nucleic acid;
; DESCRIPTION: Synthetic DNA oligonucleotide
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 399..902
; US-08-318-193-9

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Best Local Similarity 86.3%; Pred. No. 1e-60;
Matches 351; Conservative 0; Mismatches 53; Indels 0; Gaps 0;

Qy 35 CCGGCATCCAGGCGCGCATCGCGCCAGCGCGCAGCCCGAGCCCGTCCACCCAGCGCGTGGG 94
Db 502 CCGCTCCGGGGCGTCTGCAGCCCCCGCGCGTCCGCTCGCGTCCAGCCAGCGTGGG 561

Qy 95 AGCAGTGAAAGCGATCCAGAGCGCGCAGGCTCTCAACCTTCCCGCGACACCGCGG 154
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Db 802 CGTCGTGCGCACCCAGATCATACGTTTCAGTTCGTTTCAGAGGAACTCAAGGACTTCC 861

Qy 395 TCCTCGTGATCCCGTTCGATCTGTTGGAGCGCGTGCAGGAGTGA 438
Db 862 TCCTCGTGATCCCGTTCGATCTGTTGGAGCGCGTGCAGGAGTGA 905

RESULT 4
US-08-318-193-1
; Sequence 1, Application US/08318193
; Patent No. 5641663
; GENERAL INFORMATION:
; APPLICANT: GARVIN, Robert T.
; APPLICANT: MALEK, Lawrence T.
; TITLE OF INVENTION: AN EXPRESSION SYSTEM FOR THE SECRETION
; OF BIOACTIVE HUMAN GRANULOCYTE MACROPHAGE COLONY
; STIMULATING FACTOR (GM-CSF) AND OTHER HETEROLOGOUS
; PROTEINS FROM STREPTOMYCES
; NUMBER OF SEQUENCES: 91

```

```

RESULT 6
5200327-2
; Patent NO. 5200327
; APPLICANT: GARVIN, ROBERT T.; MALEK, LAWRENCE T.
; TITLE OF INVENTION: EXPRESSION SYSTEM FOR THE SECRETION OF
; BIOACTIVE HUMAN GRANULOCYTE MACROPHAGE COLONY STIMULATING
; FACTOR (GM-CSF) AND OTHER HETEROLOGOUS PROTEINS FROM
; STREPTOMYCES
; NUMBER OF SEQUENCES: 24
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/224,568
; FILING DATE: 26-JUL-1988
; SEQ ID NO:2
; LENGTH: 896
5200327-2

Query Match 69.5%; Score 318.4; DB 6; Length 896;
Best Local Similarity 89.3%; Pred. NO. 1.5e-60;
Matches 343; Conservative 0; Mismatches 41; Indels 0; Gaps 0;

QY 55 GGCCTACGCGGACGCCGACGCCGTCACCCACGCCGTGGGAGCAGCTGAACGCCGATCCAG 114
db 513 GCCCGCGCGGTGCGCCCTCGCCGTGCACCCAGCCGTGGGAGACGCTCAACGCCGATCCAG 572

```

QY	115	GAGGCCCGAGGCTCCTCAACCTCTCCCGGAGACCCGCCCGGAGATGAACGAGACCGTG	174
Db	573	GAGGCCCGCGCTGCTCAACCTCTCGGGGAGACCGCGCCGAGATGAACGAGACCGTG	632
QY	175	GAGGTGATCTCCGAGATGTTGATCTCCAGAGACCGACTGCCCTCCAGACCCCGCCTCGAG	234
Db	633	GAGGTGATCTCGAGATGTTGATCTCGAGAGCCACGTGCCCTCCAGACCCGCCCTCGAG	692
QY	235	CTGTACAAGCAGGGCTCTCCGGCGGAGCCTCAACAAGCTCAAGGCCCGCTCACCATGATG	294
Db	693	CTGTACAAGCAGGGCTCTCCGGGCGAGCCTCAACAACTCAAGGGCGCTGACCATGATG	752
QY	295	GGGTCCCACTACAAGCAGCACTGCCACCGACCCCGGAGACCTCTCTGGCCACCCGATGC	354
Db	753	GGGTCCCACTACAAGCACTGCCCGCCCAACCGCGGAGACGTGCTGCGCCACCCGATGC	812
QY	355	ATCACCTTTCGAGAGCTTCAAGGAGAACTCAAGGACTTCCTCTCGTGATCCCGTTCGAC	414
Db	813	ATCACGTTTCGAGTCGTTCAAGAGNACTGAAGACTTCCTCTCGTGATCCCGTTCGAC	872
QY	415	TCCTGGAGCCGGTCGAGAGTGA	438
Db	873	TCCTGGAGCCGGTCGAGAGTGA	896

RESULT 7  
 US-08-318-193-7  
 ; Sequence 7, Application US/08318193  
 ; Patent No. 5641663  
 ; GENERAL INFORMATION:  
 ; APPLICANT: GARVIN, Robert T.  
 ; APPLICANT: MALEK, Lawrence T.  
 ; TITLE OF INVENTION: AN EXPRESSION SYSTEM FOR THE SECRETION  
 ; TITLE OF INVENTION: OF BIOACTIVE HUMAN GRANULOCYTE MACROPHAGE COLONY  
 ; TITLE OF INVENTION: STIMULATING FACTOR (GM-CSF) AND OTHER HETEROLOGOUS  
 ; TITLE OF INVENTION: PROTEINS FROM STREPTOMYCES  
 ; NUMBER OF SEQUENCES: 91  
 ; CORRESPONDENCE ADDRESS:  
 ; ADDRESSEE: Foley & Lardner  
 ; STREET: 1800 Diagonal Road, Suite 500  
 ; CITY: Alexandria  
 ; STATE: Virginia  
 ; COUNTRY: USA  
 ; ZIP: 22313-0299  
 ; COMPUTER READABLE FORM:  
 ; MEDIUM TYPE: Floppy disk  
 ; COMPUTER: IBM PC compatible  
 ; OPERATING SYSTEM: PC-DOS/MS-DOS  
 ; SOFTWARE: Patent in Release #1.0, Version #1.25  
 ; CURRENT APPLICATION DATA:  
 ; APPLICATION NUMBER: US/08/318,193  
 ; FILING DATE:  
 ; CLASSIFICATION: 435  
 ; PRIOR APPLICATION DATA:  
 ; APPLICATION NUMBER: US/07/935,314  
 ; FILING DATE:  
 ; APPLICATION NUMBER: US 07/224,568  
 ; ATTORNEY/AGENT INFORMATION:  
 ; NAME: BENT, Stephen A.  
 ; REGISTRATION NUMBER: 29,768  
 ; REFERENCE/DOCKET NUMBER: 18740/116 CACO  
 ; TELECOMMUNICATION INFORMATION:  
 ; TELEPHONE: (703)836-9300  
 ; TELEFAX: (703)683-4109  
 ; TELEX: 899149  
 ; INFORMATION FOR SEQ ID NO: 7:  
 ; SEQUENCE CHARACTERISTICS:  
 ; LENGTH: 900 base pairs  
 ; TYPE: nucleic acid  
 ; STRANDEDNESS: double  
 ; TOPOLOGY: linear  
 ; MOLECULE TYPE: Other nucleic acid;

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; DESCRIPTION: Synthetic DNA oligonucleotide
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 399..893
; US-08-318-193-7

Query Match      69.5%; Score 318.4; DB 1; Length 900;
Best Local Similarity 89.3%; Pred. No. 1.6e-60;
Matches 343; Conservative 0; Mismatches 41; Indels 0; Gaps 0;

QY      55  GCGCCAGCGCGCAGCCGCGCTCCACCCAGCCGTGGGAGCAGCGTGAACGGATCCAG 114
DB      513  GCCCCGCCCGTCTGCGCCCTCGCCGTCAGCCAGCCGTGGGAGCAGCGTCAACGGATCCAG 572

QY      115  GAGGCCCGCAGGCTCTCTAACTCTCCCGCGACACCGCCGCGAGATGAACGAGACCGGTG 174
DB      573  GAGGCCCGCGCTGCTCAACTCTCTCGCGGACACGCGCCGCGAGATGAACGAGACCGGTG 632

QY      175  GAGGTGATCTCGAGATGTTTGGATCTTCAGGAGCCGACCTGCTCTCAGACCCGCGCTCGAG 233
DB      633  GAGGTGATCTCGAGATGTTTGGATCTTCAGGAGCCGACCTGCTCTCAGACCCGCGCTCGAG 692

QY      235  CTGTACAAGCAGCGGCTCCGCGCAGCCTCACCAAGCTCAAGGCGCCGCTCACCATGATG 294
DB      693  CTGTACAAGCAGCGGCTCCGCGGCGAGCTCACCAAGCTCAAGGCGCGCTGACCATGATG 752

QY      295  GCGTCCCACTACAAGCAGACATGCCCCACGACCCCGGAGACCTCTCGCGCACCCAGATC 354
DB      753  GCGTCCCACTACAAACAGACACTGCCCCCCACGCGGAGAGCGTCTGCGCACCCAGATC 812

QY      355  ATCAGCTTCGAGAGCTTCAAGGAGAACCTCAAGGACTTCTCTCTCTGATGATCCCGTTGAC 414
DB      813  ATCAGCTTCGAGTCGTTCAAGGAGAACCTCAAGGACTTCTCTCTCTGATGATCCCGTTGAC 872

QY      415  TGCTGGAGCCGCTGCGAGGTGA 438
DB      873  TGCTGGAGCCGCTGCGAGGTGA 896

RESULT 8
5200327-1
; Patent No. 5200327
; APPLICANT: GARVIN, ROBERT T.; MALEK, LAWRENCE T.
; TITLE OF INVENTION: EXPRESSION SYSTEM FOR THE SECRETION OF
; BIOACTIVE HUMAN GRANULOCYTE MACROPHAGE COLONY STIMULATING
; FACTOR (GM-CSF) AND OTHER HETEROLOGOUS PROTEINS FROM
; STREPTOMYCES
; NUMBER OF SEQUENCES: 24
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/224,568
; FILING DATE: 26-JUL-1998
; SEQ ID NO:1:
; LENGTH: 386
5200327-1

Query Match      68.8%; Score 315; DB 6; Length 386;
Best Local Similarity 91.7%; Pred. No. 7.5e-60;
Matches 333; Conservative 0; Mismatches 30; Indels 0; Gaps 0;

QY      76  CCGTCAACCAGCCGTGGGAGCACTGTAACGCGATCCAGGAGCCCGCAGCTCCTCAAC 135
DB      24  CCGTCAACCAGCCGTGGGAGCACTCAACGCGATCCAGGAGCCCGCAGCTCCTCAAC 83

QY      136  CTCTCCCGGACACCGCCGCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTC 195
DB      84  CTCTCCCGGACACCGCCGCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTC 143

QY      196  GATCTCCAGGAGCCGACCTGCTCTCCAGACCCGCGCTCGAGCTGTACAAGCAGGGGCTCCGC 255
DB      144  GACTTCAGGAGCCGACGTCCTCCAGACCCGCGCTCGAGCTGTACAAGCAGGGGCTCCGC 203

QY      256  GGCAGGCTCACCAAGTCTAAGGGCGCGCTCACCATGATGGGCTCCCACTACAAGCAGCAC 315

```

Db 204 GGCAGCCTCACCAGCTCAAGGGCGCTGACCATGATGGCTCCCACTACAAACAGCAC 263  
QY 316 TGCCACCGACCCCGGAGACTCTCTGGCCACCCAGATACATCACTTCGAGAGCTTCAAG 375  
Db 264 TGCCCCCACCACGCGGAGAGCTGTGGCCACCCAGATATCACTACGTTGAGTGTTCAG 323  
QY 376 GAGAACTCAAGAGACTTCTCTCTGATCCCGTTCGACTGCTGGAGCGGTGCAAGGAG 435  
Db 324 GAGAACTGAAGAGACTTCTCTCTGATCCCTTCGACTGCTGGAGCGGTGCAAGGAG 383  
QY 436 TGA 438  
Db 384 TGA 386

RESULT 9  
US-10-188-056-31  
; Patent No. 5200327  
; APPLICANT: GARVIN, ROBERT T.; MALEK, LAWRENCE T.  
; TITLE OF INVENTION: EXPRESSION SYSTEM FOR THE SECRETION OF  
; BIOACTIVE HUMAN GRANULOCYTE MACROPHAGE COLONY STIMULATING  
; FACTOR (GM-CSF) AND OTHER HETEROLOGOUS PROTEINS FROM  
; STREPTOMYCES  
; NUMBER OF SEQUENCES: 24  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/07/224,568  
; FILING DATE: 26-JUL-1988  
; SEQ ID NO:1:  
; LENGTH: 386

Query Match 68.8%; Score 315; DB 6; Length 386;  
Best Local Similarity 91.7%; Pred. No. 7.5e-60;  
Matches 333; Conservative 0; Mismatches 30; Indels 0; Gaps 0;

QY 76 CCGTCCACCCAGCGGTGGGAGCAGCTGAAGCGCATCCAGAGCGCCGCGAGGCTTCTCAAC 135  
Db 24 CCGTCCACCCAGCGGTGGGAGCAGCTGAAGCGCATCCAGAGCGCCGCGAGGCTTCTCAAC 83  
QY 136 CTCTCCCGGACACCGCGCGGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTTC 195  
Db 84 CTCTCCGGGACACCGCGCGGAGATGAACGAGACCGTGGAGGTGATCTCGGAGATGTTTC 143  
QY 196 GATCTCCAGAGCGGACCTGCTCTCCAGACCGCGCTCGAGCTGTACAAGCAGGCGCTCCGC 255  
Db 144 GACTTCAGAGCGGCGGCTGCTCTCCAGACCGCGCTCGAGCTGTACAAGCAGGCGCTCCGC 203  
QY 256 GGCAGGCTACCAAGCTCAAGGCGCGCTCACCATGATGGCGTCCCACTACAAGAGGAC 315  
Db 204 GGCAGGCTACCAAGCTCAAGGCGCGCTGACCATGATGGCGTCCCACTACAAGAGGAC 263  
QY 316 TGCCACCGACCCCGGAGACTCTCTGGCCACCCAGATATCACTTCGAGAGCTTCAAG 375  
Db 264 TGCCCCCACCACGCGGAGAGCTGTGGCCACCCAGATATCACTACGTTGAGTGTTCAG 323  
QY 376 GAGAACTCAAGAGACTTCTCTCTGATCCCGTTCGACTGCTGGAGCGGTGCAAGGAG 435  
Db 324 GAGAACTGAAGAGACTTCTCTCTGATCCCTTCGACTGCTGGAGCGGTGCAAGGAG 383  
QY 436 TGA 438  
Db 384 TGA 386

RESULT 10  
US-10-188-056-31  
; Sequence 31, Application US/10188056  
; Patent No. 6809191  
; GENERAL INFORMATION:  
; APPLICANT: Qiu, Jian-Tai  
; APPLICANT: Lai, Wan-Ching  
; APPLICANT: Chu, Yong Liang  
; APPLICANT: Li, Frank Q.

; TITLE OF INVENTION: Improved GM-CSF Nucleic Acid Sequences  
; FILE REFERENCE: 3781-004-27  
; CURRENT APPLICATION NUMBER: US/10/188,056  
; CURRENT FILING DATE: 2002-09-26  
; NUMBER OF SEQ ID NOS: 34  
; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 31  
; LENGTH: 435  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-188-056-31

Query Match 68.3%; Score 312.6; DB 4; Length 435;  
Best Local Similarity 82.8%; Pred. No. 2.6e-59;  
Matches 357; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

QY 8 GGATGCACCACACACACCACTCTCCGGCATCGAGGCGCGCATGGGCCGCGCA 67  
Db 5 GGCTGCAGAGCTGTCTCTGTGGGCACCGTGGCCTGCGAGCATCAGGCTCCCGCCAGAA 64  
QY 68 GCCGAGCCCGTCCACCCAGCCGTGGGAGCACGTGAACGCGATCCAGGAGCGCCGAGGC 127  
Db 65 GCCCAGCCCTCCACCCAGCCCTGGGAGCACGTGAACGCGATCCAGGAGCGCCAGAGGC 124  
QY 128 TCCTCAACCTCTCCCGGACACCGCCGCGAGATGAACGAGACCGTGGAGGTGATCTCCG 187  
Db 125 TGCTGAACCTGTCCAGAGACACCGCCGCGAGATGAACGAGACCGTGGAGGTGATCAGCG 184  
QY 188 AGATGTTTCATCTCCAGGAGCGGACCTGCTCCAGACCGCCCTCGAGCTGTACAAGCAGG 247  
Db 185 AGATGTTTCATCTCCAGGAGCGGACCTGCTCCAGACCGCCCTCGAGCTGTACAAGCAGG 244  
QY 248 GCTCCCGCGGAGCTTCAACAGCTCAAGGCGCGCTCAACATGATGGCGTCCCACTACA 307  
Db 245 GACTCGGGGAGCTTCAACAGCTCAAGGAGCGGCTGATGATGGCGGAGCTTCAACAGG 304  
QY 308 AGAGACATGCGCCACCGACCCGCGAGACCTCTCGCCGACCCAGATCATCCTTCGAGA 367  
Db 305 AGCAGCACTGCGCTTCCACACCGAGACGAGCTGCGCCACCCAGATCATCCTTCGAGA 364  
QY 368 GCTTCAAGAGAGACCTCAAGGACTTCTCTCGTGTATCCGTTCCGACTGCTGGAGCGCG 427  
Db 365 GCTTCAAGAGAGACCTCAAGGACTTCTCTCGTGTATCCGTTCCGACTGCTGGAGCGCG 424  
QY 428 TGCAGGAGTGA 438  
Db 425 TGCAGGAGTGA 435

RESULT 11  
US-10-188-056-33  
; Sequence 33, Application US/10188056  
; Patent No. 6809191  
; GENERAL INFORMATION:  
; APPLICANT: Qiu, Jian-Tai  
; APPLICANT: Lai, Wan-Ching  
; APPLICANT: Chu, Yong Liang  
; APPLICANT: Li, Frank Q.  
; TITLE OF INVENTION: Improved GM-CSF Nucleic Acid Sequences  
; FILE REFERENCE: 3781-004-27  
; CURRENT APPLICATION NUMBER: US/10/188,056  
; CURRENT FILING DATE: 2002-09-26  
; NUMBER OF SEQ ID NOS: 34  
; SOFTWARE: FastSEQ for Windows Version 4.0  
; SEQ ID NO 33  
; LENGTH: 435  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-188-056-33

Query Match 67.9%; Score 311; DB 4; Length 435;  
Best Local Similarity 82.6%; Pred. No. 5.7e-59;  
Matches 356; Conservative 0; Mismatches 75; Indels 0; Gaps 0;

QY 8 GGATGACACCAACCAACCACTCTCTCCGGCATCGAGGGCCGCGATGGCGCCAGCGCGCA 67  
DB 5 GGCTGCAGAGCCTGCTCTCTGGGACCGTGGCATCGAGCATCAGCGCTCCCGCCAGAA 64  
QY 68 GCCGAGCCGCTCCACCCAGCGCTGGAGCAGTGAACCGGATCCAGGAGGCCCGCAGGC 127  
DB 65 GCCCCAGCCCTCCACCCAGCCCTGGAGCAGTGAACCGGATCCAGGAGGCCCGCAGGC 124  
QY 128 TCTCAACCTCTCCCGCGACACCGCCGCGAGATGAACGAGACCGTGGAGGTGATCTCCG 187  
DB 125 TGCTGACCTGTCAGAGACACCGCCGCGAGATGAACGAGACCGTGGAGGTGATCAGGC 184  
QY 188 AGATGTCGATCTCCAGGAGCGACCTGCTCCAGACCCGCTCCAGCTGTGAACGAGG 247  
DB 185 AGATGTCGACCTGAGGAGCCACCTGCTCCAGACCCGCTCCAGCTGTGAACGAGG 244  
QY 248 GCTCCGCGGAGCCTCACAAGCTCAAGGCGCGCTCAACCATGATGGCGTCCACTACA 307  
DB 245 GACTCGGGGAGCCTGACCAAGCTGAAGGAGCCCGCTGACCATGATGGCCAGCCACTACA 304  
QY 308 AGCAGACTGCCACCGACCGGAGACCTCTCGGCGCCAGACCTCAACATCATCCTTCGAGA 367  
DB 305 AGCAGACTGCCCTCCACACCGGAGACCTGCTGCGCCAGACCTCAACATCATCCTTCGAGA 364  
QY 368 GCTTCAAGGAGAACTCAAGGACTTCTCTCTCGTATCCCGTTCGACTCTGGAGCCGG 427  
DB 365 GCTTCAAGGAGAACTGAAGGACTTCTCTCTGCTGATCCCTTCGACTCTGGAGCCGG 424  
QY 428 TGCAGAGTGA 438  
DB 425 TGCAGAGTGA 435

## RESULT 12

US-08-469-318-55  
; Sequence 55, Application US/08469318  
; Patent No. 6022535  
; GENERAL INFORMATION:  
; APPLICANT: Multivariant IL-3 Hematopoiesis Fusion  
; TITLE OF INVENTION: Protein  
; NUMBER OF SEQUENCES: 196  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/469,318  
; FILING DATE:  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/446,872  
; FILING DATE:  
; INFORMATION FOR SEQ ID NO: 55:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 777 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
US-08-469-318-55

Query Match 51.9%; Score 283.6; DB 3; Length 777;  
Best Local Similarity 81.8%; Pred. No. 5.8e-53;  
Matches 328; Conservative 0; Mismatches 74; Indels 0; Gaps 0;  
QY 34 TCCGGCATCGAGGGCGCGATGGCGGCGAGCGCGAGCCCGGATCCACCGCGGTGG 93  
DB 376 TCTGGCGGGCTCCACATGCGAGCGGTCTGTTCCCGGCTCCCGCTACCGAGCGGTGG 435  
QY 94 GAGCAGTGAAACGCGATCCAGGAGGCCCGCAGGCTCTCAACCTCTCCCGCGACACCGCC 153

DB 436 GAACACGTGAATGCCATCCAGAGGCCCGCGCTCTCTGAACCTGAGTAGACACTGCT 495  
QY 154 GCCGAGATGAACGAGACCGTGGAGGTGATCTCTCCGAGATGTTGATCTCCAGAGCCGACC 213  
DB 496 GCTGAGATGAATGAACAGTAGAAGTATATCAGAAATGTTTGACCTCCAGGAGCGACT 555  
QY 214 TGCCTCCAGACCCGCTCGAGCTGTACAGCAGGCGCTCCGCGGAGCCTCAACGAGCTC 273  
DB 556 TGCCTACAGACCCGCTCGAGCTGTACAGCAGGCGCTCCGCGGAGCCTCAACGAGCTC 615  
QY 274 RAGGCGCCGCTCACCATGATGGCTCCCACTACAGCAGCAGTCCCAACCGCCGCGAG 333  
DB 616 AAGGCGCCCTTGACCATGATGGCCAGCCACTACAAGCAGCACTGCCCTCAACCCCGGAA 675  
QY 334 ACCTCTCGCGCCACCCAGATCATCATCTTCGAGAGCTTCAAGGAGAACTCAAGGACTTC 393  
DB 676 ACTTCTGTGCAACCCAGATTCACCTTTGAAAGTTTCAAGAGAACTCAAGGACTTC 735  
QY 394 CTCTCTCGTATCCCTTCGACTCTGGGAGCCGCTGCGAGGAG 435  
DB 736 CTGCTTGTATCCCTTTGACTCTGCTGGAGCCAGTCCAGGAG 777

## RESULT 13

US-08-468-609A-55  
; Sequence 55, Application US/08468609A  
; Patent No. 6030812  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; APPLICANT: Bauer, S. C.  
; APPLICANT: Braford-Goldberg, Sarah R.  
; APPLICANT: Caparon, Mairé H.  
; APPLICANT: Easton, Alan M.  
; APPLICANT: Klein, Barbara K.  
; APPLICANT: McKearn, John P.  
; APPLICANT: Olins, Peter O.  
; APPLICANT: Paik, Kuman  
; APPLICANT: Thomas, John W.  
; TITLE OF INVENTION: Fusion Proteins Comprising Multiply Mutated Interleukin-3 (IL-3)  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
; STREET: P. O. Box 5110  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60680  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/468,609A  
; FILING DATE: 06-JUN-1995  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Bennett, Dennis A.  
; REGISTRATION NUMBER: 34,547  
; REFERENCE/DOCKET NUMBER: C-2790/3  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (314)737-6986  
; TELEFAX: (314)737-6972  
; INFORMATION FOR SEQ ID NO: 55:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 777 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double



```

; APPLICATION NUMBER: US 08/192,325
; FILING DATE: 14-FEB-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Bennett, Dennis A.
; REGISTRATION NUMBER: 34,547
; REFERENCE/DOCKET NUMBER: C-2790/1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)737-6986
; TELEFAX: (314)737-6972
; INFORMATION FOR SEQ ID NO: 55:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 777 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-446-872A-55

Query Match          61.9%; Score 283.6; DB 3; Length 777;
Best Local Similarity 81.6%; Pred. No. 5.8e-53;
Matches 328; Conservative 0; Mismatches 74; Indels 0; Gaps 0

QY 34 TC CGGCATCAGGGCCGCATGGCCAGCGCGCAGCCCGAGCCCGTCCACCCAGCGGTGG 93
DB 376 TCTGGCGCGCGCTCCAACTATGCGACCGCGCTCGTTCCCGCTCCCGCTACCCAGCGGTGG 435

QY 94 GAGCAGTGAACCGGATCCAGGAGGCGCGCAGGCTCTCTCAACCTCTCCCGGACACCGCC 153
DB 436 GAACAGTGTAATGCCATCCAGGAGGCGCGCGCTCTCTGAACTGTAGTAGAGACACTGCT 495

QY 154 GCCGAGATGAACGAGACCGGTGGAGGTGATCTCCGAGATGTTTCGATCTCCAGGAGCCGACC 213
DB 496 GCTGAGATGAATGAACAGTAGAAGTGATATCAGAAATGTTTGNCTCTCAGGAGCCGACT 555

QY 214 TGCCTCAGACCCGCTCTGAGCTGTGAACAGAGGGCTCTCCGCGCAGCTCTACCAAGCTC 273
DB 556 TGCCTACAGACCCGCTTGGAGCTGTGAACAGAGGGCTCTCGGGGCGACTCACCAAGCTC 615

QY 274 AAGGGCCGCTCACCATGATGGCGTCCCACTACAGCAGCACTTGCCCAACCGACCCCGGAG 333
DB 616 AAGGGCCCCCTTGACCATGATGGCCAGCCACTAAGCAGCACTTGCCCTCCAACCCCGGNA 675

QY 334 ACCTCTCGGCCACCCAGATCATCAGCTTCGAGAGCTTCAAGGAGAACTTCAAGGACTTC 393
DB 676 ACTTCTGTGCAACCCAGATATATCACTTTGAAGTTTCAAGAGAACTTGAAGGACTTC 735

QY 394 CTCCTCGTATCCGTTGCACTGCTGGAGCCCGGTGCAGAG 435
DB 736 CTCCTGTGATCCCTTTGACTGCTGGAGCCAGTTCAGAG 777

```

```

RESULT 15
US-08-762-227A-55
; Sequence 55, Application US/08762227A
; Patent No. 6436387
; GENERAL INFORMATION:
; APPLICANT: Abrams, Mark A.
; Bauer, S. C.
; Braford-Goldberg, Sarah R.
; Caparon, Mairé H.
; Easton, Alan M.
; Klein, Barbara K.
; McKearn, John P.
; Oline, Peter O.
; Paik, Kuman
; Thomas, John W.
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis
; Fusion Protein
; NUMBER OF SEQUENCES: 197
; CORRESPONDENCE ADDRESS:
; ADDRESSSEE: Dennis A. Bennett, G.D. Searle & Co
; Corporate Patent Dept.
; STREET: P. O. Box 5110

```

CITY: Chicago  
STATE: Illinois  
COUNTRY: USA  
ZIP: 60680  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/762,227A  
FILING DATE: 09-Dec-1996  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/192,325  
FILING DATE: 14-FEB-1994  
APPLICATION NUMBER: US 08/446,872  
FILING DATE: 06-JUN-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: Bennett, Dennis A.  
REGISTRATION NUMBER: 34,547  
REFERENCE/DOCKET NUMBER: C-2790/5  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (708)470-6501  
TELEFAX: (708)470-6881  
INFORMATION FOR SEQ ID NO: 55:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 777 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
SEQUENCE DESCRIPTION: SEQ ID NO: 55:  
US-08-762-227A-55

Query Match 61.9%; Score 283.6; DB 3; Length 777;  
Best Local Similarity 81.6%; Pred. No. 5.8e-53;  
Matches 328; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

QY	34	TCGGGATCGAGGGCGCGATGGCGCGCAGCGCGAGCGCGCTCCACCGAGCGGTGG	93
Db	376	TCGGGGGGGGCTTCAACATGGCACCAGGCTGTTCCTCCCGTCCCGCTACCCAGCGGTGG	435
QY	94	GAGCAGTGAAACCGGATCCAGGAGGCGCGAGGCTCTCAACCTCTCCCGCGACACCGCC	153
Db	436	GAACACGTGAATGCCATCCAGGAGGCGCGCGCTCTCTGAACTGTAGTAGAGACACTGCT	495
QY	154	GCCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTTGGATCTCCAGGAGCGGACC	213
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QY	274	AAGGGCCCGCTCACCATGTATGGCTCCCACTACAAGCAGCACTGCCACCGACCCCGGAG	333
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QY	334	ACCTCTGCGCCACCCGATCATCAGCTTCGAGAGCTTCAAGGAGAACCTCAAGGACTTC	393
Db	676	ACTTCTGTGCAACCCAGATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTC	735
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Search completed: March 11, 2005, 17:26:43  
Job time : 148 secs

GenCore version 5.1.6  
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 11, 2005, 15:19:06 ; Search time 2523 Seconds  
(without alignments)  
6909.804 Million cell updates/sec

Title: US-10-723-083-1  
Perfect score: 458  
Sequence: 1 cggccggatgaccaccca.....getaggtgcagcatgcgcg 458

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479089

Minimum DB seq length: 0  
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Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : EST.\*

1: gb\_est1.\*  
2: gb\_est2.\*  
3: gb\_hic.\*  
4: gb\_est3.\*  
5: gb\_est4.\*  
6: gb\_est5.\*  
7: gb\_est6.\*  
8: gb\_gssi.\*  
9: gb\_gssi2.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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5	276.8	60.4	695	2	BE669962 7e27g08.x
6	276.8	60.4	895	2	BE873976 601484045
7	276.2	60.3	658	5	BX111836 BX111836
8	273.6	59.7	592	7	CF341802 TGEStzvJ4
9	273.6	59.7	592	7	CF370966 TGEStzvJ5
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11	264.8	57.8	718	6	CD369973 UI-H-FT1-
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13	256.4	56.0	565	2	BF938995 7r03f11.x
14	243.6	53.2	666	6	CA307828 UI-H-FT1-
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17	208.6	45.5	561	6	CB457551 714908 MA
18	207.2	45.2	672	7	CF614774 CES009198
19	202.4	44.2	511	6	CB430266 606148 MA
20	200.2	43.7	608	6	CD367244 UI-H-FT2-
21	197.6	43.1	572	4	BM539160 hb05e10.9
22	191.2	41.7	336	2	AW951121 EST363191
23	176.6	38.6	549	1	AI677936 wc88f12.x
24	175.2	38.3	423	2	AW784714 zb77g08.9

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C	36	67	14.6	1138	2	BE636680
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C	39	59	12.9	160	7	CF341980
C	40	59	12.9	925	9	CNS0091P
C	41	58.2	12.7	830	9	CM010233
C	42	58	12.7	136	7	CF341168
C	43	57.8	12.6	881	9	CG339578
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## ALIGNMENTS

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ACCESSION AW207707  
VERSION AW207707.1 GI:6507203  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1 (bases 1 to 588)  
AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
TITLES National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
Tumor Gene Index  
JOURNAL Unpublished (1997)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: [cgapps@mail.nih.gov](mailto:cgapps@mail.nih.gov)  
Oligo-dT track not found, Not 1 site shown in beginning of sequence  
is likely internal to the message. cDNA library preparation: M.B.  
Scores Lab Clone distribution: NCI-CGAP clone distribution  
Information can be found through the I.M.A.G.E. Consortium/LLNL at:  
[www-bio.llnl.gov/bbrp/image/image.html](http://www-bio.llnl.gov/bbrp/image/image.html)  
Seq primer: M13 Forward  
POLYA=No.

FEATURES  
source

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NCI CGAP Sub4 library is a subtracted library derived from  
the NCI CGAP Sub2 library which is a subtracted library  
derived from the NCI CGAP Sub1 library, which is a  
subtracted library derived from BI. BI constitutes a  
mixture of 21 normalized or subtracted NCI\_CGAP  
libraries: NCI\_CGAP\_Co4, NCI\_CGAP\_Pr22, NCI\_CGAP\_Pr28,  
NCI\_CGAP\_Co10, NCI\_CGAP\_Co16, NCI\_CGAP\_Kid5,  
NCI\_CGAP\_Kid12, NCI\_CGAP\_Kid3, NCI\_CGAP\_Kid1,  
NCI\_CGAP\_Lym2, NCI\_CGAP\_Br2, NCI\_CGAP\_Co8, NCI\_CGAP\_CLL1,  
NCI\_CGAP\_Le12, NCI\_CGAP\_Br23, NCI\_CGAP\_Lu5,  
NCI\_CGAP\_Lu24, NCI\_CGAP\_Lu19, NCI\_CGAP\_GC4, NCI\_CGAP\_GC6,

NCI CGAP Brn25. These 21 libraries were pooled and a single-stranded DNA preparation of the resulting mixture was used as a tracer in a subtractive hybridization with a driver whose composition is detailed below:

NCI CGAP Kid3 pool 1 : LLAM 3334-3337, 3682-3683, 3798-3803 (IMAGE Clonoids 1322376-1323911, 1456008-1456775, 1500552-1502855) NCI CGAP Kid5 pool 1 : LLAM 3338-3342, 3722-3725, 3776-3778 (IMAGE Clonoids 1323912-1325831, 1471368-1472903, 1492104-1493255)

NCI CGAP Lu5 pool 1 : LLAM 3575-3582, 3851-3854 (IMAGE Clonoids 144920-1417991, 1520904-1522439) NCI CGAP GC4 pool 1 : LLAM 3164-3167, 3716-3720, 3733-3735 (IMAGE Clonoids 1257096-1258631, 1469064-1470983, 1475592-1476743) NCI CGAP Pr22 pool 1 : LLAM 2457-2459, 2758-2759, 3062-3068 (IMAGE Clonoids 985608-986759, 1101192-1101959, 1217928-1220615) NCI CGAP Co10 pool 1 : LLAM 2644-2653, 2871-2872 (IMAGE Clonoids 1057416-1061255, 1144584-1145351) Subtraction was performed as previously described [Bonaldo, Lennon & Soares (1996): Normalization and Subtraction: Two Approaches To Facilitate Gene Discovery. Genome Research 6, 791-806.]

TAG\_TISSUE=colon  
TAG\_LIB=NCI CGAP\_Co4  
TAG\_SEQ=CTTCG"

## ORIGIN

Query Match 60.4%; Score 276.8; DB 2; Length 588;  
Best Local Similarity 77.5%; Pred. No. 7e-50;  
Matches 335; Conservative 0; Mismatches 97; Indels 0; Gaps 0;

QY 8 GGATGCACCAACCACCACTCTCCGGCATCGAGGCGGCATGGCGCCAGCGCGA 67  
DB 41 GGCTGCAGAGCCTGCTCTTGGGACATGTGGCTTCGACATCTCTGCACCGCCGCT 100

QY 68 GCCGAGCCCGTCCACCGACCGCTGGGAGACGTGAACGGGATCCAGAGGCCCGCAGGC 127  
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QY 128 TCCTCAACCTCTCCGCGACCGCCGAGATGAACGAGCCGCGGAGGTGATCTCG 187  
DB 161 TCCTGAACCTGAGTAGAGACATGCTGCTGAGATGAATGAACAGTAGAAGTATCTCAG 220

QY 188 AGATGTTTCGATCTCCAGGAGCCGACCTGCTCCAGACCGCCCTCGAGCTGTACAAGCAG 247  
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QY 248 GCCTCGCGGCGAGCTTACCAAGCTCAAGGCGCGCTCACCATGATGGCGTCCCACTACA 307  
DB 281 GCCTGGGGGAGCCTTCAAGCTCAAGGGGCCCTTGACCATGATGGCCAGCCACTACA 340

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QY 368 GCTTCAAGGAGAACCTCAAGGACTTCTCCTCGTGATCCCGTTTCGACTGCTGGGAGCCGG 427  
DB 401 GTTTCGAAGAGAACCTCAAGGACTTCTCCTCGTGATCCCGTTTCGACTGCTGGGAGCCAG 460

QY 428 TCAGGAGTGAG 439  
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## RESULT 2

BE218982 660 bp mRNA linear EST 03-JUL-2000  
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FACTOR PRECURSOR (HUMAN); mRNA sequence.

ACCESSION BE218982  
VERSION BE218982.1 GI:8906300  
KEYWORDS EST.  
SOURCE Homo sapiens (human)

## ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 660)

NCI CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: [cgapbs@mail.nih.gov](mailto:cgapbs@mail.nih.gov)

Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.

Emmert-Buck, M.D., Ph.D.

CDNA Library Preparation: M. Bento Soares, Ph.D.

DNA Sequencing Arrayed by: Greg Lennon, Ph.D.

Clone distribution: NCI-CGAP clone distribution information can be

found at [image.llnl.gov](http://image.llnl.gov)

Seq primer: -40UP from Gibco

High quality sequence stop: 445.

Location/Qualifiers

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/note="Organ: lung; Vector: p773D-Pac (Pharmacia) with a

modified polylinker; Plasmid DNA from the normalized

library NCI CGAP Lu5 was prepared, and ss circles were

made in vitro. Following HAP purification, this DNA was

used as tracer in a subtractive hybridization reaction.

The driver was PCR-amplified cDNAs from a pool of 5,000

clones made from the same library (clonoids

1414920-1417991 and 1520904-1522439). Subtraction by Bento

Soares and M. Fatima Bonaldo. "

"

"

"

"

Query Match 60.4%; Score 276.8; DB 2; Length 660;

Best Local Similarity 77.5%; Pred. No. 7e-50;

Matches 335; Conservative 0; Mismatches 97; Indels 0; Gaps 0;

QY 8 GGATGCACCAACCACCACTCTCCGGCATCGAGGCGGCATGGCGCCAGCGCGA 67

DB 9 GGCTGCAGAGCCTGCTCTTGGGACATGTGGCTTCGACATCTCTGCACCGCCGCT 68

QY 68 GCCGAGCCCGTCCACCGACCGCTGGGAGCACGTGAACCGCATCCAGGAGGCCCGCAGGC 127

DB 69 CGCCAGCCCCAGCAGCGCCCTGGGAGCATGTGAATGCCATCCAGGAGGCCCGCGTC 128

QY 128 TCCTCAACCTCTCCCGCGACACCGCGCGAGATGAACGAGACCGCTGGAGGTGATCTCG 187

DB 129 TCCTGAACCTGATGATAGACATCTGCTGCTGAGATGAATGAACAGTAGAAGTATCTCAG 188

QY 188 AGATGTTTCGATCTCCAGGAGCGGACCTGCTCCAGACCGCCCTCGAGCTGTACAAGCAGG 247

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QY 248 GCCTCGCGGCGAGCCTTCAACAGCTCAAGGGGCCCGCTCAACATGATGGCGTCCCACTACA 307

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QY 308 AGCAGCACTGCCACCGACCGCCGAGACCTCTCGCGCCACCCAGATCATCACCCTTCGAGA 367

DB 309 AGCAGCACTGCCCTTCAACCCCGGAACCTTCTGTGTCAACCCCAAGATTATCACCTTTGAAA 368

QY 368 GCTTCAAGGAGAACCTCAAGGACTTCTCCTCGTGATCCCGTTTCGACTGCTGGGAGCCGG 427

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FACTOR PRECURSOR (HUMAN);, mRNA sequence.
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AI912784
VERSION
AI912784.1 GI:5632639
KEYWORDS
EST.
SOURCE
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ORGANISM
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 666)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
TITLE
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
CDNA Library Arrayed by: Greg Lemmon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
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library NCI CGAP Lu5 was prepared, and ss circles were
made in vitro. Following HAP purification, this DNA was
used as tracer in a subtractive hybridization reaction.
The driver was PCR-amplified cDNAs from a pool of 5,000
clones made from the same library (clonoids
1414920-1417991 and 1520904-1522439). Subtraction by Bento
Soares and M. Fatima Bonaldo."
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Best Local Similarity 77.5%; Pred. No. 7e-50;
Matches 335; Conservative 0; Mismatches 97; Indels 0; Gaps 0;
QY 8 GGATGACACACACACACACCTCTCTCGGATCGAGGCCGATCGCGCGCGCA 67
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Db 9 GGCTGAGAGAGCTGCTCTTGGGACACTGTGGCCCTGCAGCATCTCTGCACCCGCGCT 68
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Qy		248	GCCTCCCGGGGAGCCTCACCAAGCTCAAGGGCCCGCTCACCATGATGGCGTCCCACTACA	307	
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Qy		308	AGCAGCATGCCCCACGACCCCGGAGACCTCGTGGGCCACCCAGATCATACACTTTCGAGA	367	
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modified polylinker; Plasmid DNA from the normalized library NCI CGAP Lu5 was prepared, and ss circles were made *in vitro*. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (clonoids 141420-1417991 and 1520304-1522439). Subtraction by Bento Soares and M. Fatima Bonaldo. "

ORIGIN

Query Match	60.4%;	Score	276.8;	DB	2;	Length	695;	
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QY	368	GCTTCAAGGAGAACCTCAAGGACTTCTCTCTCGTGAATCCCGTTTCGACTGCTGGGACCGG	427					
DB	405	GTTTCAAGAGAACCTGAAGGACTTCTCTGTCATCCCTTTGACTGCTGGAGCCAG	464					
QY	428	TGCAGGAGTGAG	439					
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VERSION	BE873976.1	GI:10322752			
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ORGANISM	Homo sapiens				
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
REFERENCE	1 (bases 1 to 895)				
AUTHORS	NIH-MGC <a href="http://mgc.nci.nih.gov/">http://mgc.nci.nih.gov/</a> .				
TITLE	National Institutes of Health, Mammalian Gene Collection (MGC)				
JOURNAL	Unpublished (1999)				
COMMENT	Contact: Robert Strausberg, Ph.D. Email: <a href="mailto:cgapbs@mail.nih.gov">cgapbs@mail.nih.gov</a> Tissue Procurement: DCTD/DTF/Gazdar CDNA Library Preparation: Life Technologies, Inc. CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Incyte Genomics, Inc. Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <a href="http://image.llnl.gov">http://image.llnl.gov</a> Plate: LLAM9663 row: j column: 12				

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      /notes="Organ: lung; Vector: pCMV-SPORT6; Site 1: NotI; Site 2: SalI; Cloned unidirectionally. Primer: Oligo dr. Average insert size 1.1 kb. Library constructed by Life Technologies."
ORIGIN
  Query Match      60.4%; Score 276.8; DB 2; Length 895;
  Best Local Similarity 77.5%; Pred. No. 7.1e-50;
  Matches 335; Conservative 0; Mismatches 97; Indels 0; Gaps 0;
QY 8 GGATGACACACACACACACACCTCTCCGGCATCGAGGCCCGCATGGCCGACGGCGCA 67
DB 3 GGCTGACAGACCTGCTGCTTGGGCACCTGGGCTGCAGCATCTCTGCACCCGCCGCT 62
QY 68 GCCGAGCCGTCACCCAGCCGTGGAGCAGCTGAACGGATCCAGAGGCCCGCAGGC 127
DB 63 CGCCAGCCCGCAGCAGCAGCCCTGGGAGCATGTGAATCCATCCAGGAGGCCGGGCTC 122
QY 128 TCCTCAACCTCTCCCGCGACACCGCCGAGATGAACGAGACCGTGGAGGTATCTCCG 187
DB 123 TCCTGAACCTGATAGAGACACTGCTGTGAGATGATGAACAGTAGATCATCTCAG 182
QY 188 AGATGTTGATTCAGAGACCGACCTGCTCCAGACCCGCCCTCGAGCTGTATAAGCAGG 247
DB 183 AAATGTTTGAACCTCCAGAGCCGACCTGCTTACAGACCCGCCCTGGAGCTGTACAAGCAGG 242
QY 248 GCCTCCGCGCAGCCTCACAAGCTCAAGGCGCGCTCACCATGATGGGTCCCATACA 307
DB 243 GCCTCGGGGCGAGCCTCACAAGCTCAAGGCGCGCTTGCATGATGGCGCAGCCTACA 302
QY 308 AGCAGCACTGCCACACGACCCCGAGACCTCTCGCGCCACCCAGATCATCACCTTCGAGA 367
DB 303 AGCAGCACTGCCCTCCAAACCCCGGAACTTCCTGTGCAACCCAGATATACCTTTGAA 362
QY 368 GCTTCAAGGAGACCTCAAGGACTTCTCTCTGTGATTCCTGTTGAGTGGAGCCGG 427
DB 363 GTTTCGAAGAGAACCTCAAGGACTTCTCTGTGATTCCTTGTCTGCTGCTGAGGAGCAG 422
QY 428 TGCAGGAGTGAG 439
DB 423 TCCAGGAGTGAG 434
RESULT 7
BX111836/c 658 bp mRNA linear EST 07-FEB-2003
LOCUS
DEFINITION
  BX111836 NCI CGAP Lu5 Homo sapiens CDNA clone IMAGp998C104061;
  IMAGE:1601601, mRNA sequence.
ACCESSION
  BX111836
VERSION
  BX111836.1 GI:27837278
KEYWORDS
  EST.
SOURCE
  Homo sapiens (human)
  ORGANISM
    Homo sapiens
    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
    Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  REFERENCE
    1 (bases 1 to 658)
    Ebert,L., Heil,O., Hennig,S., Neubert,P., Partsch,E., Peters,M.,
    Radelof,U., Schneider,D. and Korn,B.
    Human Unigeneset - RZPD3
    JOURNAL
    Unpublished (2003)
    COMMENT
    Contact: Ina Rolfs
    RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
    Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany
```

```
RZPD; IMAGp998C104061.
RZPDLIB; I.M.A.G.E. CDNA Clone Collection;
Human Unigeneset - RZPD3 (RZPDLIB No.972)
http://www.rzpd.de/CloneCards/cgi-
bin/showLib.pl.cgi/response?libNo=972 Contact: Ina Rolfs
RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
Heubnerweg 6, D-14059 Berlin, Germany
Tel: +49 30 32639 101
Fax: +49 30 32639 111
www.rzpd.de
This clone is available royalty-free from RZPD;
contact RZPD (clone@rzpd.de) for further information. Seq primer:
M13r, Primer sequence: TTTACACAGGAACAGCTATGAC.
FEATURES
  Location/Qualifiers
    1..658
      /organism="Homo sapiens"
      /mol_type="mRNA"
      /db_xref="taxon:9606"
      /clone_lib="IMAGp998C104061 ; IMAGE:1601601"
      /tissue_type="carcinoid"
      /lab_host="DHI0B"
      /clone_lib="NCI CGAP Lu5"
      /notes="Organ: lung; Vector: pT7T3D-Pac (Pharmacia) with a
      modified polylinker; 1st strand cDNA was prepared from
      neuroendocrine lung carcinoid, and was then primed with a
      Not I - oligo(dT) primer. Double-stranded cDNA was ligated
      to Eco RI adaptors (Pharmacia), digested with Not I and
      cloned into the Not I and Eco RI sites of the modified
      pT7T3 vector. Library is normalized. Library was
      constructed by Bento Soares and M. Fatima Bonaldo."
ORIGIN
  Query Match      60.3%; Score 276.2; DB 5; Length 658;
  Best Local Similarity 82.3%; Pred. No. 9.5e-50;
  Matches 317; Conservative 0; Mismatches 68; Indels 0; Gaps 0;
QY 55 GCGCCAGCGGCGAGCCGCTCCACCCAGCCGCTGGGAGCAGCTGACGCGATCCAG 114
DB 619 GCACCCGCCGCTCGCCCCAGCCCGACGACGCGCTGGGAGCATGTGAATGCCATCCAG 560
QY 115 GAGGCCGCGAGGCTCTCTCAACCTCTCCCGCAGACCCGCGCCGAGATGAACGAGACCGTG 174
DB 559 GAGGCCGCGGCTCTCTGAACTGTAGTAGACACTGCTGTGAGATGAATGAACAGTA 500
QY 175 GAGGTGATCTCCGAGATGTTGATGATTCAGGAGCCGACCTGCTCTCAGACCCGCCCTCGAG 234
DB 499 GAAGTCATCTCAGAAATGTTTGACCTCCAGGAGCCGACCTGCTCAGACCCGCCCTGGAG 440
QY 235 CTGTACAAGCGGGCTCTCGCGGCGAGCTCCACCAAGCTCAAGGGCCGCTCACCATGATG 294
DB 439 CTGTACAAGCGGGCTCTCGCGGCGAGCTCCACCAAGCTCAAGGGCCGCTTGAACCATGATG 380
QY 295 GCGTCCCACTACAAGCAGCACTGCCCCAGCCGAGACCTCTCTGCGCCACCCAGATC 354
DB 379 GCCAGCACTACAAGCAGCACTGCCCTCCAAACCCCGGAACTTCTCTGTGCAACCCAGATT 320
QY 355 ATCACCTTCCAGAGCTTCAAGGAGAACCTCAAGGACTTCTCTCTGTGATCCGCTTCGAC 414
DB 319 ATCACCTTGAAGAGTTTCAAGAGAACCTTGAAGGACTTCTCTGCTGTGTCATCCCTTTGAC 260
QY 415 TGCTGGAGCCGCTGCAGGAGTGAG 439
DB 259 TGCTGGAGCCAGTCCAGGAGTGAG 235
RESULT 8
CF341802 584 bp mRNA linear EST 18-AUG-2003
LOCUS
DEFINITION
  TgESTzyj43f02.y1 Tg CAST Tachyzoite cDNA Library Toxoplasma gondii
  CDNA clone TgESTzyj43f02.y1 5' similar to SW:CSF2 HUMAN P04141
  GRANULOCYTE-MACROPHAGE COLONY-STIMULATING FACTOR PRECURSOR ;, mRNA
  sequences.
ACCESSION
  CF341802
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VERSION      CF341802.1  GI:33831915
KEYWORDS     EST.
SOURCE       Toxoplasma gondii
ORGANISM     Toxoplasma gondii
REFERENCE    1 (bases 1 to 584)
AUTHORS      Tang,K., Cole,R., Fogarty,S., Sibley,L.D., Ajioaka,J.A., White,M.,
              Clifton,S., Pape,D., Martin,J., Wyllie,T., Dante,M., Marra,M.,
              Hillier,L., Kucaba,T., Theising,B., Bowers,Y., Gibbons,M.,
              Ritter,E., Bennett,J., Franklin,C., Tsagareishvili,R., Ronko,I.,
              Kennedy,S., Maguire,L., Waterston,R. and Wilson,R.
TITLE        Toxoplasma EST Project
JOURNAL      Unpublished (2001)
COMMENT      Contact: Clifton, S.
              Toxoplasma EST Project
              Washington University School of Medicine
              4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
              Tel: 314 286 1800
              Fax: 314 286 1810
              Email: toxo@watson.wustl.edu
              Contact David Sibley (toxos@borcim.wustl.edu) for further
              information relating to organism, libraries, or clone availability.
              Seq primer: -40UP from Gibco.
              Location/Qualifiers
                1..584
                  /organism="Toxoplasma gondii"
                  /mol_type="mRNA"
                  /db_xref="taxon:5811"
                  /clone="TgESTzyj43f02.y1"
                  /dev_stage="Tachyzoite"
                  /lab_host="Electroten Blue cells (Stratagene)"
                  /clone_lib="Tg CAST Tachyzoite cDNA Library"
                  /notes="Vector: Modified pBluescript (pBS SK+); Site 1:
                  BamHI; Site 2: EcoRI; The cDNA library was constructed by
                  Keliang Tang, and Robert Cole at Washington University.
                  cDNA was synthesized from poly(A)+ mRNA using the
                  template-switching PCR method (SMART cDNA Kit, BD
                  Biosciences). First strand cDNA was reverse transcribed
                  using the CDS III/3' primer and a 5' template switch
                  primer (Smart IV primer). The product of the first strand
                  synthesis was PCR amplified using the same primer set and
                  the fragments were digested with SfiI. The fragments were
                  size selected, ligated into a modified pBluescript vector
                  (obtained from Michael White, Montana State University)
                  containing directional SfiI sites, and electroporated into
                  ElectroTen Blue cells. Vector: SfiI sites were added to
                  the multiple cloning region of pBluescript SK+ between the
                  BamHI/EcoRI sites. The modified polylinker has the
                  following sequence: 5'GAATTGGCCATTACGGCC(G)n-- insert--
                  GCGCGCTCGCCACGGATCC3' where n=3-4 G nucleotides.
                  WARNING: the library contains a small percentage of cDNAs
                  derived from the human host cells. Library materials
                  provided by David Sibley, Washington University."

FEATURES     source
SOURCE       CF370966
LOCUS        CF370966.1  GI:34318212
DEFINITION   Toxoplasma gondii
KEYWORDS     Eukaryota; Alveolata; Apicomplexa; Coccidia; Eimeriida;
SOURCE       Sarcocystidae; Toxoplasma.
ORGANISM     1 (bases 1 to 592)
REFERENCE    Tang,K., Cole,R., Fogarty,S., Sibley,L.D., Ajioaka,J.A., White,M.,
              Clifton,S., Pape,D., Martin,J., Wyllie,T., Dante,M., Marra,M.,
              Hillier,L., Kucaba,T., Theising,B., Bowers,Y., Gibbons,M.,
              Ritter,E., Bennett,J., Franklin,C., Tsagareishvili,R., Ronko,I.,
              Kennedy,S., Maguire,L., Waterston,R. and Wilson,R.
AUTHORS      Kennedy,S., Maguire,L., Waterston,R. and Wilson,R.
TITLE        Toxoplasma EST Project
JOURNAL      Unpublished (2001)
COMMENT      Contact: Clifton, S.
              Toxoplasma EST Project
              Washington University School of Medicine
              4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
              Tel: 314 286 1800
              Fax: 314 286 1810
              Email: toxo@watson.wustl.edu
              Contact David Sibley (toxos@borcim.wustl.edu) for further
              information relating to organism, libraries, or clone availability.
              Seq primer: -40UP from Gibco.
              Location/Qualifiers
                1..592
                  /organism="Toxoplasma gondii"
                  /mol_type="mRNA"
                  /db_xref="taxon:5811"
                  /clone="TgESTzyj58e12.y1"
                  /dev_stage="Tachyzoite"
                  /lab_host="Electroten Blue cells (Stratagene)"
                  /clone_lib="Tg CAST Tachyzoite cDNA Library"
                  /notes="Vector: Modified pBluescript (pBS SK+); Site 1:
                  BamHI; Site 2: EcoRI; The cDNA library was constructed by
                  Keliang Tang, and Robert Cole at Washington University.
                  cDNA was synthesized from poly(A)+ mRNA using the
                  template-switching PCR method (SMART cDNA Kit, BD
                  Biosciences). First strand cDNA was reverse transcribed
                  using the CDS III/3' primer and a 5' template switch
                  primer (Smart IV primer). The product of the first strand
                  synthesis was PCR amplified using the same primer set and
                  the fragments were digested with SfiI. The fragments were
                  size selected, ligated into a modified pBluescript vector
                  (obtained from Michael White, Montana State University)
                  containing directional SfiI sites, and electroporated into
                  ElectroTen Blue cells. Vector: SfiI sites were added to
                  the multiple cloning region of pBluescript SK+ between the
                  BamHI/EcoRI sites. The modified polylinker has the
                  following sequence: 5'GAATTGGCCATTACGGCC(G)n-- insert--
                  GCGCGCTCGCCACGGATCC3' where n=3-4 G nucleotides.
                  WARNING: the library contains a small percentage of cDNAs
                  derived from the human host cells. Library materials
                  provided by David Sibley, Washington University."

ORIGIN
Query Match      60.1%; Score 275.2; DB 7; Length 584;
Best Local Similarity 77.3%; Pred. No. 1.6e-49;
Matches 334; Conservative 0; Mismatches 98; Indels 0; Gaps 0;

QY 8  GGATGCACCAACCAACCACTCTCTCGGCATCGAGGGCCGATCGGCGCCAGCGCA 67
DB 37 GGCTGCAGAGCCCTGCTCTTGGGCACTGTGGCTTCGACATCTCTGCACCCGCCGCT 96
QY 68 GCCGAGCCGCTCACCCAGCCCTGGGAGCACTGACGCGATCCAGGAGCCCGCAGCG 127
DB 97 CGCCACGAGCCCGAGCAGCAGCCCTGGGAGCATGTGATGATCCATCCAGGAGCCCGCGTC 156
QY 128 TCCTCAACTCTCTCCCGGACACCGCCCGGAGATGAACAGAGACCGCTGGAGGTATCTCCG 187
DB 157 TCCTGAACCTGATAGAGACACTGCTGCTGAGATGATGATGAACAGTAGATCATCTCAG 216
QY 188 AGATGTTGATCTCCAGGAGCCGACCTGCTCCAGACCCCGCTTCGAGCTGTACAGAGG 247

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```
RESULT 14
CA307828/c
LOCUS
DEFINITION
  UI-H-FTI-bhx-f-10-0-UI.s1 NCI CGAP_FTI Homo sapiens cDNA clone
  UI-H-FTI-bhx-f-10-0-UI 3', mRNA sequence.
ACCESSION
CA307828
VERSION
CA307828.1 GI:24470882
SOURCE
  Homo sapiens (human)
ORGANISM
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  1 (bases 1 to 666)
  NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
  National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
  Tumor Gene Index
  Unpublished (1997)
  Contact: Robert Strausberg, Ph.D.
  Email: cgaps-r@mail.nih.gov
  cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
  DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
  Clone Distribution: Clone distribution information can be obtained
  from Dr. M. Bento Soares, bento-soares@uiowa.edu
  The following repetitive elements were found in this cDNA
  sequence: 1-50, >AT richLow_complexity 64-133,
  >(TAAA)n#Simple_repeat
  Seq primer: M13_FORWARD
  POLYA=Yes.

FEATURES
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    1..666
    /organism="Homo sapiens"
    /mol_type="mRNA"
    /db_xref="taxon:9606"
    /clone="UI-H-FTI-bhx-f-10-0-UI"
    /tissue_type="Alveolar Macrophage"
    /dev_stage="Adult"
    /lab_host="DH10B (Life Technologies)"
    /clone_lib="NCI CGAP FTL"
    /note="Organ: Lung; Vector: p773-Pac (Pharmacia) with a
    modified polylinker; Site 1: EcoR I; Site 2: Not I;
    NCI CGAP_FTI is a normalized cDNA library constructed from
    a pool of 81 RNA samples from Alveolar Macrophages
    challenged with different treatments. The mRNA samples
    were a mixture of these conditions (times refer to
    incubations following isolation by bronchoalveolar lavage)
    (some normal donor macrophages were cultured in some of
    the conditions, other donor macrophages in different
    conditions). The mRNA samples were pooled for library
    construction. Control 0 hours; Control 3 hours; Control 24
    hours; LPS 100 ng/ml, 3 hours; LPS 100 ng/ml, 24 hours;
    PMA 10 ng/ml, 3 hours; PMA 10 ng/ml, 24 hours; Klebsiella
    moi 10, 3 hours; Klebsiella moi 10, 24 hours; Staph aureus
    moi 10, 3 hours; Staph aureus moi 10, 24 hours; Adenoviral
    vector (Ad5 CMV eGFP), moi 500, 3 hours; Adenoviral vector
    (Ad5 CMV eGFP), moi 500, 24 hours; wt adenovirus moi 500,
    3 hours; wt adenovirus moi 500, 24 hours; Ad vector + LPS
    3 hours; Ad vector + LPS 24 hours; wt adenovirus + LPS 3
    hours; wt adenovirus + LPS 24 hours. The library was
    normalized according to Bonaldo, Lennon and Soares, Genome
    Research, 6:791-806, 1996. First strand cDNA synthesis was
    primed with an oligo-dT primer containing a Not I site.
    Double stranded cDNA was ligated to an EcoR I adaptor,
    digested with Not I, and cloned directionally into
    p773-Pac vector. The oligonucleotide used to prime the
    synthesis of first-strand cDNA contains a library tag
    sequence that is located between the Not I site and the
    (dT)18 tail. The sequence tag for this library is
    GGCCATGCCG. The tissue was provided by Dr. Gary W.
    Hunninghake of the University of Iowa.
    TAG_TISSUE=Human Lung Alveolar Macrophage
    TAG_LIB=UI-H-FTI

RESULT 15
CD368851/c
LOCUS
DEFINITION
  UI-H-FTI-bjx-m-02-0-UI.s1 NCI CGAP_FTI Homo sapiens cDNA clone
  UI-H-FTI-bjx-m-02-0-UI 3', mRNA sequence.
ACCESSION
CD368851
VERSION
CD368851.1 GI:31152941
KEYWORDS
  EST.
SOURCE
  Homo sapiens (human)
ORGANISM
  Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  1 (bases 1 to 661)
  NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
  National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
  Tumor Gene Index
  Unpublished (1997)
  Contact: Robert Strausberg, Ph.D.
  Email: cgaps-r@mail.nih.gov
  Tissue procurement: Dr. Gary W. Hunninghake, U of I
  cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
  DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
  Clone Distribution: Distribution information can be found at
  http://genome.uiowa.edu/distribution/cgap.html
  The following repetitive elements were found in this cDNA
  sequence: 65-134, >(TAAA)n#Simple_repeat
  Seq primer: M13_FORWARD
  POLYA=Yes.

FEATURES
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    /tissue_type="Alveolar Macrophage"
    /dev_stage="Adult"
    /lab_host="DH10B (Life Technologies)"
    /clone_lib="NCI CGAP_FTI"
    /note="Organ: Lung; Vector: p773-Pac (Pharmacia) with a
    modified polylinker; Site 1: EcoR I; Site 2: Not I;
    modified polylinker; Site 1: EcoR I; Site 2: Not I;
    TAG_TISSUE=Human Lung Alveolar Macrophage
    TAG_LIB=UI-H-FTI

TAG_SEQ=GGCCATGCCG"
ORIGIN
  Query Match 53.2%; Score 243.6; DB 6; Length 666;
  Best Local Similarity 83.4%; Pred. No. 1.1e-42;
  Matches 276; Conservative 0; Mismatches 55; Indels 0; Gaps 0;
  QY 109 ATCCAGGAGGCCGCGAGGCTCTCTCAACCTCTCCCGGACACACCGCCCGAGATGAACGAG 168
  |||||
  DB 665 ATCCAGGAGGCCGCGGCTCTCTGAACTGAGTAGAGACACTGCTGCTGAGATGAATGAA 606
  |||||
  QY 169 ACCGTGGAGGTGATCTCCGAGATGTTGATCTCCAGGAGCGGACCTGCTCTCAGACCCGC 228
  |||||
  DB 605 ACAGTAGAAGTCACTCTCAGAAATGTTTGACCTCCAGGAGCGGACCTGCTCTCAGACCCGC 546
  |||||
  QY 229 CTCGAGCTCTACAAGCAGGCGCTCCGCGGCGAGCTCACCAGCTCAAGGCGCGCTCACC 288
  |||||
  DB 545 TTGGAGCTGTACAAGCAGGCGCTCGCGGCGAGCTCACCAGCTCAAGGCGCGCTTGCACC 486
  |||||
  QY 289 ATGATGGCTCTCCACTTACAAAGCAGCAGCTGCCACCGACCCCGGAGACCTCTCTCGCCACC 348
  |||||
  DB 485 ATGATGGCCAGCCANTACAAGCAGCAGCTGCCCTCCAAACCCCGGAAACTTCTGTGCAACC 426
  |||||
  QY 349 CAGATCATCAGCTTCGAGAGCTTCAAGGAGAACCTCAAGGAGCTTCTCTCTCGTATCCCG 408
  |||||
  DB 425 CAGATTATCACCTTTGAAAGTTTCAAAGAGAACCTGAGGAGCTTTCTGCTGTGTCATCCC 366
  |||||
  QY 409 TTGCACTGCTGGAGCGCGTGCAGGAGTGAG 439
  |||||
  DB 365 TTGCACTGCTGGAGCGCGTGCAGGAGTGAG 335
  |||||

FEATURES
  source
    1..661
    /organism="Homo sapiens"
    /mol_type="mRNA"
    /db_xref="taxon:9606"
    /clone="UI-H-FTI-bjx-m-02-0-UI"
    /tissue_type="Alveolar Macrophage"
    /dev_stage="Adult"
    /lab_host="DH10B (Life Technologies)"
    /clone_lib="NCI CGAP_FTI"
    /note="Organ: Lung; Vector: p773-Pac (Pharmacia) with a
    modified polylinker; Site 1: EcoR I; Site 2: Not I;
    modified polylinker; Site 1: EcoR I; Site 2: Not I;
    TAG_TISSUE=Human Lung Alveolar Macrophage
    TAG_LIB=UI-H-FTI
```

NCI\_CGAP\_FTL1 is a normalized cDNA library constructed from a pool of 81 RNA samples from Alveolar Macrophages challenged with different treatments. The mRNA samples were a mixture of these conditions (times refer to incubations following isolation by bronchoalveolar lavage) (some normal donor macrophages were cultured in some of the conditions, other donor macrophages in different conditions). The mRNA samples were pooled for library construction. Control 0 hours; control 3 hours; control 24 hours; LPS 100 ng/ml, 3 hours; LPS 100 ng/ml, 24 hours; PMA 10 ng/ml, 3 hours; PMA 10 ng/ml, 24 hours; Klebsiella moi 10, 3 hours; Klebsiella moi 10, 24 hours; Staph aureus moi 10, 3 hours; Staph aureus moi 10, 24 hours; Adenoviral vector (Ad5 CMV egfp), moi 500, 3 hours; Adenoviral vector (Ad5 CMV egfp), moi 500, 24 hours; wt adenovirus moi 500, 3 hours; wt adenovirus moi 500, 24 hours; Ad vector + LPS 3 hours; Ad vector + LPS 24 hours; wt adenovirus + LPS 3 hours; wt adenovirus + LPS 24 hours. The library was normalized according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pT73-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is GGCCATGCCG. The tissue was provided by Dr. Gary W. Hunninghake of the University of Iowa.

TAG\_TISSUE=Human Lung Alveolar Macrophage  
TAG\_LIB=UI-H-FTL1  
TAG\_SEQ=GGCCATGCCG"

## ORIGIN

Query Match	50.4%	Score 230.8	DB 6	Length 661
Best Local Similarity	83.5%	Pred. No. 6.7e-40		
Matches 273	Conservative 0	Mismatches 53	Indels 1	Gaps 1
Qy 113	AGGAGGCCGCGAGGCTCTCAACCTTCGCGGACACCGCGCGGAGATGAACGAGACCG 172			
Db 661	AGGAGGCCGCGGCTCTCTGAACTGAGTAGAGACACTGCTGCTGAGATGAATGAACAG 602			
Qy 173	TGGAGGTGATCTCGAGATGTTGATCTCCAGGAGCGGACCTGCTCCAGACCGGCTCG 232			
Db 601	TGAAGTCACTCAGAAATG-TNGACCTCCAGGAGCGGACCTGCTCCACAGACCGGCTCG 543			
Qy 233	AGCTGTACAAGCAGGCGCTCCGCGGAGCTCACCAGCTCAAGGCGCGCTCACCCTGA 292			
Db 542	AGCTGTACAAGCAGGCGCTTCGCGGAGAGCTCACCAGCTCAAGGCGCGCTTGAACATGA 483			
Qy 293	TGGCGTCCCACTACAAGCAGCACTGCCCCACCGACCCCGGAGACCTCTCGGCCACCCAGA 352			
Db 482	TGCCAGGCCACTACAAGCAGCACTGCCCCCAACCCCGGAACTTCTGTGCAACCCAGA 423			
Qy 353	TCATCACTTCGAGAGTTCAAGAGAACCTCAAGACTTCCTCTCGTGATCCCGTTGG 412			
Db 422	TTATCACTTTGAAGTTTCAAGAGAACCTGAAGGACTTTCTGCTTGTGATCCCTTTG 363			
Qy 413	ACTGCTGGGAGCGGTGAGAGTGAG 439			
Db 362	ACTGCTGGGAGCGGTGAGAGTGAG 336			

Search completed: March 11, 2005, 17:24:17  
Job time : 2532 secs



GenCore version 5.1.6  
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 11, 2005, 09:51:36 ; Search time 420 Seconds  
(without alignments)  
6455.337 Million cell updates/sec

Title: US-10-723-083-1

Perfect score: 458

Sequence: 1 cggccggatgcaccacca.....gtagcgtagcagcatgccg 458

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N\_Geneseq\_16Dec04:\*

1: Geneseqn1980s:\*

2: Geneseqn1990s:\*

3: Geneseqn2000s:\*

4: Geneseqn2001as:\*

5: Geneseqn2001bs:\*

6: Geneseqn2002as:\*

7: Geneseqn2002bs:\*

8: Geneseqn2003as:\*

9: Geneseqn2003bs:\*

10: Geneseqn2003cs:\*

11: Geneseqn2003ds:\*

12: Geneseqn2004as:\*

13: Geneseqn2004bs:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	319.2	69.7	905	2	AAQ03221 DNA fragm
2	318.4	69.5	896	2	AAQ03220 DNA fragm
3	315	68.8	386	2	AAQ03219 DNA fragm
4	312.6	68.3	435	12	AD120788 Human GM-
5	311	67.9	435	12	AD120790 Human GM-
6	284.4	62.1	435	12	AD076023 Human GM-
7	283.6	61.9	777	2	AAQ97169 pMON13022
8	283.6	61.9	777	3	AAA03723 Human int
9	283.6	61.9	777	6	ABX00012 Human int
10	283.6	61.9	777	12	ADJ14267 DNA relat
11	283	61.8	402	2	AAQ97208 pMON13012
12	283	61.8	402	3	AAA03771 Human G-C
13	283	61.8	402	6	ABX00086 Human int
14	283	61.8	402	12	ADJ14388 DNA relat
15	281	61.4	822	2	AAQ97183 pMON13035
16	281	61.4	822	3	AAA03737 Human int
17	281	61.4	822	6	ABX00026 Human int
18	281	61.4	822	12	ADJ14281 DNA relat
19	281	61.4	903	2	AAQ97180 pMON13031
20	281	61.4	903	3	AAA03734 Human int

21	281	61.4	903	6	ABX00023 Human int
22	281	61.4	903	12	ADJ14278 DNA relat
23	277.8	60.7	1538	10	ADF31962 Human GM-
24	277.8	60.7	1538	10	ADF31989 Human gra
25	277.6	60.6	415	1	AAAN90383 Synthetic
26	277.6	60.6	415	1	AAAN90274 Synthetic
27	277	60.5	2211	12	ADL16719 Human stu
28	276.8	60.4	448	12	ADL16729 Human gra
29	276.8	60.4	763	2	AAQ04018 Granulocy
30	276.8	60.4	781	12	ADN07714 Human G-C
31	276.8	60.4	781	12	ADP10387 Reference
32	276.8	60.4	787	1	AAAN60364 Human gra
33	276.8	60.4	787	2	AAQ84865 Clone pCD
34	276.8	60.4	789	3	AAA35017 Human ade
35	276.8	60.4	789	3	AAF21139 Human low
36	276.8	60.4	789	4	AAH28217 Nucleotid
37	276.8	60.4	789	10	ABZ96833 Human nuc
38	276.8	60.4	789	11	ABD20682 Human pul
39	276.8	60.4	2385	2	AAAT72725 Her2-GM-C
40	276.8	60.4	5115	3	AAA35020 Human ade
41	276.8	60.4	5115	10	ABZ96836 Human nuc
42	276.8	60.4	5115	11	ABD20685 Human pul
43	276.8	60.4	5115	11	ABD20685 Human pul
44	276.8	60.4	9629	2	AAAT14600 pXJCL-hGM
45	276.2	60.3	1011	8	ABX63509 Human CDN

#### ALIGNMENTS

##### RESULT 1

AAQ03221  
ID AAQ03221 standard; DNA; 905 BP.

XX AAQ03221;

DT 12-JUL-1990 (first entry)

XX DNA fragment of pABO-GMCSF encodes granulocyte macrophage colony

DE stimulating factor (GM-CSF).

XX Granulocyte macrophage; colony stimulating factor; GM-CSF; cancer; da.

OS Streptomyces sp.

XX EP352707-A.

XX 31-JAN-1990.

XX 24-JUL-1989; 89EP-00113607.

XX 25-JUL-1988; 88CA-00572956.

XX (CANG-) CANGENE CORP.

XX Garvin RT, Malek LT;

XX WPI; 1990-031296/05.

XX gene expression system directing secretion of protein in Streptomyces -

PT contg. structural gene, esp. for granulocyte macrophage colony

PT stimulating factor, and regulatory sequence.

XX Disclosure; Fig 4; 39pp; English.

XX GM-CSF, or a wide range of other proteins, is secreted in unglycosylated

CC form, with correctly positioned intramolecular disulphide bonds and full

CC biological activity. pAEO.GMCSF contains the BamHI HindIII fragment with

CC an aminoglycoside phosphotransferase promoter and the protease B-endo H

XX hybrid signal peptide. GM-CSF is potentially useful in cancer treatment

XX Sequence 905 BP; 131 A; 367 C; 298 G; 109 T; 0 U; 0 Other;

```
Query Match      69.7%; Score 319.2; DB 2; Length 905;
Best Local Similarity 86.9%; Pred. No. 5.8e-49;
Matches 351; Conservative 0; Mismatches 53; Indels 0; Gaps 0;

QY 35 CCGGATCAGGGCCGATGGCCAGCGCGAGCCCGGAGCCCGTCCACCGAGCGGTGGG 94
DB 502 CCGCCTCCGGGGGCTGCGAGCCCGCCCGCGTCCGCGTCCGACCCAGCGGTGGG 561
QY 95 AGCAGTGAACGGATCCAGGAGGCGCGCAGGCTCTCAACCTCTCCCGCGACACCGCGG 154
DB 562 AGCAGTGAACGGATCCAGGAGGCGCGCGTCTCAACCTCTCCCGCGACACCGCGG 621
QY 155 CCGAGATGAACGAGACCGGTGGAGGTGATCTCCGAGATGTTCCGATCTCCAGGAGCGGACCT 214
DB 622 CCGAGATGAACGAGACCGGTGGAGGTGATCTCCGAGATGTTCCGATCTCCAGGAGCGGACCT 681
QY 215 GCCTCAGACCGCCTCGAGCTGTACAAGCAGGGGCTCCGGCGCAGCTTCAACGACTCA 274
DB 682 GCCTCAGACCGCCTCGAGCTGTACAAGCAGGGGCTCCGGCGCAGCTTCAACGACTCA 741
QY 275 AGGGCCCGCTCACCATGATGGGTCCCACTACAAGCAGCACTGCCCAACCGACCCCGGAGA 334
DB 742 AGGGCCCGCTCACCATGATGGGTCCCACTACAAGCAGCACTGCCCAACCGACCCCGGAGA 801
QY 335 CCTCTGCGCCACCCAGATCATCCTTCGAGAGCTTCAAGGAGAACTCAAGGACTTCC 394
DB 802 CGTGTGCGCCACCCAGATCATCCTTCGAGTCTGTTCAAGGAGAACTCAAGGACTTCC 861
QY 395 TCCTGTGATCCGTTGCACTGCTGGAGCGCGTGCAGGAGTGA 438
DB 862 TCCTGTGATCCGTTGCACTGCTGGAGCGCGTGCAGGAGTGA 905

RESULT 2
AAQ03220
ID AAQ03220 standard; DNA; 896 BP.
AC AAQ03220;
XX
XX
DT 12-JUL-1990 (first entry)
DE DNA fragment of pAPO-GM-CSF encodes granulocyte macrophage colony
DE stimulating factor (GM-CSF).
XX
XX KW Granulocyte macrophage; colony stimulating factor; GM-CSF; cancer; ds.
XX OS Streptomyces sp.
XX PN EP352707-A.
XX PD 31-JAN-1990.
XX PF 24-JUL-1989; 89EP-00113607.
XX PR 25-JUL-1988; 88CA-00572956.
XX PA (CANG-) CANGENE CORP.
XX PI Garvin RT, Malek LT;
XX DR WPI; 1990-031296/05.
XX PT gene expression system directing secretion of protein in Streptomyces -
XX stimulating factor, and regulatory sequence.
XX PS Disclosure; Fig 3; 39pp; English.
XX CC GM-CSF, or a wide range of other proteins, is secreted in unglycosylated
XX form, with correctly positioned intramolecular disulphide bonds and full
XX biological activity. pAPO-GM-CSF contains the BamHI HindIII fragment with
XX an aminoglycoside phosphotransferase promoter and the protease B signal
XX peptide. GM-CSF is potentially useful in cancer treatment

Query Match      69.5%; Score 318.4; DB 2; Length 896;
Best Local Similarity 89.3%; Pred. No. 8.1e-49;
Matches 343; Conservative 0; Mismatches 41; Indels 0; Gaps 0;

QY 55 GCGCCAGCGCGCAGCCCGAGCCCGTCCACCCAGCCGTGGAGCAGCTGAACCGGATCCAG 114
DB 513 GCGCCCGCGCGTCCGCTCGCCGTCGACCCAGCCGTGGAGCAGCTCAACGCGATCCAG 572
QY 115 GAGGCCCGCAGGCTCTCAACCTCTCCCGCGACACCGCGCGGAGATGAACGAGACCGTG 174
DB 573 GAGGCCCGCGGCTCTCAACCTCTCCCGCGGACACCGCGCGGAGATGAACGAGACCGTG 632
QY 175 GAGGTGATCTCCGAGATGTTCCGATCTCCAGGAGCCGACCTGCTCCAGACCCGCTCCGAG 234
DB 633 GAGGTGATCTCCGAGATGTTCCGATCTCCAGGAGCCGACCTGCTCCAGACCCGCTCCGAG 692
QY 235 CTGTACAAGCAGGGGCTCCGCGGAGCTTCAAGAGTCAACAGGCTCAAGGGCCGCTCAACGATG 294
DB 693 CTGTACAAGCAGGGGCTCCGCGGAGCTTCAAGAGTCAACAGGCTCAAGGGCCGCTCAACGATG 752
QY 295 GCGTCCCACTACAAGCAGGACTGCCCAACCGACCCCGGAGACCTCTCGCGCCACCCAGATC 354
DB 753 GCGTCCCACTACAAGCAGGACTGCCCAACCGACCCCGGAGACCTCTCGCGCCACCCAGATC 812
QY 355 ATCACCTTCGAGAGCTTCAAGGAGAACTTCAAGGAGTTCCTCTCGTGATCCGTTTCGAC 414
DB 813 ATCACGTTTCGAGTCTGTTCAAGGAGAACTTCAAGGAGTTCCTCTCGTGATCCGTTTCGAC 872
QY 415 TCCTGGAGCGCGTGCAGGAGTGA 438
DB 873 TCCTGGAGCGCGTGCAGGAGTGA 896

RESULT 3
AAQ03219
ID AAQ03219 standard; DNA; 386 BP.
AC AAQ03219;
XX
XX
DT 12-JUL-1990 (first entry)
DE DNA fragment encodes granulocyte macrophage colony stimulating factor (GM
DE -CSF).
XX
XX KW Granulocyte macrophage; colony stimulating factor; GM-CSF; cancer; ds.
XX OS Streptomyces sp.
XX PN EP352707-A.
XX PD 31-JAN-1990.
XX PF 24-JUL-1989; 89EP-00113607.
XX PR 25-JUL-1988; 88CA-00572956.
XX PA (CANG-) CANGENE CORP.
XX PI Garvin RT, Malek LT;
XX DR WPI; 1990-031296/05.
XX PT gene expression system directing secretion of protein in Streptomyces -
XX stimulating factor, and regulatory sequence.
XX PS Claim 4; Fig 1; 39pp; English.
XX CC GM-CSF, or a wide range of other proteins, is secreted in unglycosylated
XX form, with correctly positioned intramolecular disulphide bonds and full
```



CC biological activity. GM-CSF is potentially useful in cancer treatment  
XX Sequence 386 BP; 73 A; 141 C; 116 G; 56 T; 0 U; 0 Other;  
Query Match 68.8%; Score 315; DB 2; Length 386;  
Best Local Similarity 91.7%; Pred. No. 3.4e-48;  
Matches 333; Conservative 0; Mismatches 30; Indels 0; Gaps 0;  
QY 76 CCGTCACCCAGCCGCGGGAGCAGCGTGAACGCGATCCAGAGGCCCGCAGGCTCTCTCAAC 135  
DB 24 CCGTCACCCAGCCGCGGGAGCAGCGTGAACGCGATCCAGAGGCCCGCAGGCTCTCTCAAC 83  
QY 136 CTCTCCCGGACACCCGCGCGGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTC 195  
DB 84 CTCTCCCGGACACCCGCGCGGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTC 143  
QY 196 GATCTCCAGGAGCCGACCTGCTCTCCAGACCCGCTCGAGCTGTACAAGCAGGCGCTCCGC 255  
DB 144 GACTTCAGGAGCCAGCGTGCCTCCAGACCCGCTCGAGCTGTACAAGCAGGCGCTCCGC 203  
QY 256 GGCAGGCTCACCAAGCTCAAGGCGCGCTCAACATGATGCGTCCCACTACAAGCAGCAC 315  
DB 204 GGCAGGCTCACCAAGCTCAAGGCGCGCTGACCATGATGCGTCCCACTACAAGCAGCAC 263  
QY 316 TGCCACCCAGCCCGGAGACCTCTGCGCCACCCAGATCATCATCTTCGAGCTTCAAG 375  
DB 264 TGCCACCCAGCCCGGAGACCTCTGCGCCACCCAGATCATCATCTTCGAGCTTCAAG 323  
QY 376 GAGAACTCAAGGACTTCTCTCTGATGCCGCTTCGACTGCTGGAGCGGTGACAGGAG 435  
DB 324 GAGAACTCAAGGACTTCTCTCTGATGCCGCTTCGACTGCTGGAGCGGTGACAGGAG 383  
QY 436 TGA 438  
DB 384 TGA 386  
RESULT 4  
AD120788  
ID AD120788 standard; DNA; 435 BP.  
XX AC AD120788;  
XX DT 22-APR-2004 (first entry)  
XX DE Human GM-CSF associated DNA #1.  
XX OS mammalian granulocyte macrophage-colony stimulating factor; GM-CSF;  
XX KW immunostimulant; Cytostatic; vaccine; ds.  
XX OS Homo sapiens.  
XX PN WO2004004742-A1.  
XX PD 15-JAN-2004.  
XX PF 03-JUL-2003; 2003WO-US020908.  
XX PR 03-JUL-2002; 2002US-00188056.  
XX PA (VAXI-) VAXIM INC.  
XX PI Qiu J, Lai W, Chu YL, Li FQ;  
XX WP; 2004-099342/10.  
XX DR New DNA sequence comprising a codon optimized sequence encoding a  
XX PT mammalian granulocyte macrophage-colony stimulating factor, useful as an  
XX PT adjuvant for enhancing an immune response of a mammal to preventive or  
XX PT therapeutic vaccine.  
XX PS Disclosure; SEQ ID NO 31; 42pp; English.  
XX XX

CC The present invention relates to a DNA sequence comprising a codon-  
CC optimized sequence encoding a mammalian granulocyte macrophage-colony  
CC stimulating factor (GM-CSF), is new, where the codon optimized sequence  
CC is different from the corresponding wild type GM-CSF encoding DNA, where  
CC a codon in the codon optimized sequence has been altered to enhance the  
CC expression of the mammalian GM-CSF. The method is useful as an adjuvant  
CC in gene therapy or in vaccination for enhancing an immune response of a  
CC mammal to a preventive or therapeutic vaccine, is also useful for  
CC treating cancer or for effecting gene therapy. Use of the DNA sequence  
CC comprising a codon optimized sequence encoding a mammalian granulocyte  
CC macrophage-colony stimulating factor as an adjuvant results in improved  
CC adjuvant activity when expressed in vivo compared with the native GM-CSF  
CC gene sequence. The present sequence represents human GM-CSF associated  
CC encoding sequence.  
XX  
SQ Sequence 435 BP; 94 A; 151 C; 128 G; 62 T; 0 U; 0 Other;  
Query Match 68.3%; Score 312.6; DB 12; Length 435;  
Best Local Similarity 82.8%; Pred. No. 9.3e-48;  
Matches 357; Conservative 0; Mismatches 74; Indels 0; Gaps 0;  
QY 8 GGATGCACCCACCCACCCACCTCTCTCCGCGATCGAGGCGCCGATGGCGCCAGCGCA 67  
DB 5 GGCTGCAGAGCTGCTCTCTCTGGGCACCGTGGCTGCAGCATCAGCGCTCCCGCCAGAA 64  
QY 68 GCGCAGAGCCGCTCCACCCAGCCGCTGGAGACGCTGAACCGGATCCAGGAGCCCGCAGGC 127  
DB 65 GCGCAGAGCCGCTCCACCCAGCCGCTGGAGACGCTGAACCGGATCCAGGAGCCCGCAGGC 124  
QY 128 TCTCAACCTCTCCCGGACACCGCGCGAGATGAACGAGACCGTGGAGGTGATCTCCG 187  
DB 125 TGTGAACCTGTCAGAGACACCGCGCGAGATGAACGAGACCGTGGAGGTGATCAGCG 184  
QY 188 AGATGTTGATCTCCAGGAGCCGACCTGCTCTCCAGACCCGCTCGAGCTGTACAAGCAGG 247  
DB 185 AGATGTTGATCTCCAGGAGCCGACCTGCTCTCCAGACCCGCTCGAGCTGTACAAGCAGG 244  
QY 248 GCTCCGCGGAGCTTCAACAGCTCAAGGCGCGCTCAACATGATGGCGTCCCACTACA 307  
DB 245 GACTCGGGGAGCTTCAACAGCTCAAGGAGCCGCTGACCATGATGGCGGAGCTTCA 304  
QY 308 AGCAGACTGCCCCACCCAGCCGCGGAGACCTCTCTCGCGCCACCCAGATCATCATCTT 367  
DB 305 AGCAGACTGCCCCACCCAGCCGCGGAGACCTGCGCGCCACCCAGATCATCATCTT 364  
QY 368 GCTTCAAGGAGAACCTCAAGGACTTCTCTCTGATCCGTTCCGACTGCTGGAGCGG 427  
DB 365 GCTTCAAGGAGAACCTCAAGGACTTCTCTCTGATCCGTTCCGACTGCTGGAGCGG 424  
QY 428 TGCAGGAGTGA 438  
DB 425 TGCAGGAGTGA 435  
RESULT 5  
AD120790  
ID AD120790 standard; DNA; 435 BP.  
XX AC AD120790;  
XX DT 22-APR-2004 (first entry)  
XX DE Human GM-CSF associated DNA #3.  
XX OS mammalian granulocyte macrophage-colony stimulating factor; GM-CSF;  
XX KW immunostimulant; Cytostatic; vaccine; ds.  
XX OS Homo sapiens.  
XX PN WO2004004742-A1.  
XX PD 15-JAN-2004.  
XX PF 03-JUL-2003; 2003WO-US020908.  
XX PR 03-JUL-2002; 2002US-00188056.  
XX PA (VAXI-) VAXIM INC.  
XX PI Qiu J, Lai W, Chu YL, Li FQ;  
XX WP; 2004-099342/10.  
XX DR New DNA sequence comprising a codon optimized sequence encoding a  
XX PT mammalian granulocyte macrophage-colony stimulating factor, useful as an  
XX PT adjuvant for enhancing an immune response of a mammal to preventive or  
XX PT therapeutic vaccine.  
XX PS Disclosure; SEQ ID NO 31; 42pp; English.  
XX XX

PF 03-JUL-2003; 2003WO-US020908.  
 XX  
 PR 03-JUL-2002; 2002US-00188056.  
 XX  
 PA (VAXI-) VAXIM INC.  
 XX  
 PI Qiu J, Lai W, Chu YL, Li FQ;  
 XX WPI; 2004-099342/10.  
 DR  
 XX New DNA sequence comprising a codon optimized sequence encoding a  
 XX mammalian granulocyte macrophage-colony stimulating factor, useful as an  
 PT adjuvant for enhancing an immune response of a mammal to preventive or  
 PT therapeutic vaccine.  
 PT  
 XX Disclosure; SEQ ID NO 33; 42pp; English.  
 PS  
 XX The present invention relates to a DNA sequence comprising a codon-  
 CC optimized sequence encoding a mammalian granulocyte macrophage-colony  
 CC stimulating factor (GM-CSF), is new, where the codon optimized sequence  
 CC is different from the corresponding wild type GM-CSF encoding DNA, where  
 CC a codon in the codon optimized sequence has been altered to enhance the  
 CC expression of the mammalian GM-CSF. The method is useful as an adjuvant  
 CC in gene therapy or in vaccination for enhancing an immune response of a  
 CC mammal to a preventive or therapeutic vaccine, is also useful for  
 CC treating cancer or for effecting gene therapy. Use of the DNA sequence  
 CC comprising a codon optimized sequence encoding a mammalian granulocyte  
 CC macrophage-colony stimulating factor as an adjuvant results in improved  
 CC adjuvant activity when expressed in vivo compared with the native GM-CSF  
 CC gene sequence. The present sequence represents human GM-CSF associated  
 CC encoding sequence.  
 XX  
 SQ Sequence 435 BP; 94 A; 152 C; 128 G; 61 T; 0 U; 0 Other;  
 Query Match 67.9%; Score 311; DB 12; Length 435;  
 Best Local Similarity 82.6%; Pred. No. 1.8e-47;  
 Matches 356; Conservative 0; Mismatches 75; Indels 0; Gaps 0;  
 QY 8 GGATGACACACACACACACACCTCTCCGGCATGAGGCGCGCATGGCGCAGCGCA 67  
 Db 5 GGCTGCAGAGCTGCTCTGCTGGGCACCTGCGCTGCAGCATCAGCGCTCCGCGCAGAA 64  
 QY 68 GCCGAGCGCGTCCACCGACCGTGGGAGCAGTGAACGGATCCAGGAGGCGCCGAGCG 127  
 Db 65 GCCCAGCGCGCTCCACCGACCGCTGGGAGCAGTGAACGCCATCCAGGAGGCGCAGCGC 124  
 QY 128 TCCTCAACCTCTCCCGGACACCGCGCGAGATGAACGAGACCGTGGAGGTGATCTCG 187  
 Db 125 TGCTGAACCTGTCCAGAGACACCGCGCGAGATGAACGAGACCGTGGAGGTGATCAGCG 184  
 QY 188 AGATGTTTCGATCTCCAGGAGCGACCTGCTCCAGACCGCGCTCCGAGCTGTACAAGCAGG 247  
 Db 185 AGATGTTTCGATCTCCAGGAGCGACCTGCTCCAGACCGCGCTCCGAGCTGTACAAGCAGG 244  
 QY 248 GCTCTCGCGGAGCTTCAACAGCTCAAGGCGCGCTCAACATGATGGCGTCCCACTACA 307  
 Db 245 GACTGGGGGAGCTTCAACAGCTCAAGGCGCGCTCAACATGATGGCGTCCCACTACA 304  
 QY 308 AGCAGCACTGCCACCGACCGCGGAGACCTCTGCGCGCACCCAGATCATCCTTCGAGA 367  
 Db 305 AGCAGCACTGCCCTCCACACCGAGACAGCTGCGCGCACCCAGATCATCCTTCGAGA 364  
 QY 368 GCTTCAAGGAGAACCTCAAGGACTTCTCTCTGATTCGCGTCTGAGTCTGGGAGCGCG 427  
 Db 365 GCTTCAAGGAGAACCTCAAGGACTTCTCTCTGATTCGCGTCTGAGTCTGGGAGCGCG 424  
 QY 428 TGCAGAGTGA 438  
 Db 425 TGCAGAGTGA 435  
 RESULT 6  
 ADQ76023

ADQ76023 standard; DNA; 435 BP.  
 ADQ76023;  
 07-OCT-2004 (first entry)  
 Human GM-CSF codon optimised coding sequence.  
 ds; gene; human; GM-CSF; codon optimisation; protein production.  
 Homo sapiens.  
 Synthetic.  
 WO2004059556-A2.  
 15-JUL-2004.  
 23-DEC-2003; 2003WO-EP014850.  
 23-DEC-2002; 2002DE-01060805.  
 (GENE-) GENEART GMBH.  
 Raab D, Graf M, Notka F, Wagner R;  
 WPI; 2004-543639/52.  
 Computer optimization of a nucleotide sequence for a protein comprises  
 evaluating test sequences with a quality function to determine the  
 optimum sequence.  
 Claim 25; SEQ ID NO 2; 83pp; German.  
 The present invention relates to a method of optimising a coding sequence  
 for expression of a protein, based on the amino acid sequence of the  
 protein. This involves the use of a computer to generate a test sequence  
 with m optimisation positions determined for a defined region, in which  
 positions the codon usage is varied. The optimum codon usage at such  
 positions is determined by means of a power function. The steps are  
 reiterated with different regions of the sequence, with the optimised  
 codons previously identified being left unchanged during subsequent  
 steps. The method can be used for expression of proteins. The present  
 sequence is a codon optimised version of the human GM-CSF coding  
 sequence.  
 Sequence 435 BP; 101 A; 132 C; 128 G; 74 T; 0 U; 0 Other;  
 Query Match 62.1%; Score 284.4; DB 12; Length 435;  
 Best Local Similarity 84.0%; Pred. No. 1.2e-42;  
 Matches 321; Conservative 0; Mismatches 61; Indels 0; Gaps 0;  
 QY 55 GCGCCAGCGCGCAGCG 114  
 Db 52 GCG 111  
 QY 115 GAGGCG 174  
 Db 112 GAGGCG 171  
 QY 175 GAGGCG 234  
 Db 172 GAGGCG 231  
 QY 235 CTGTCAAGCAGCG 294  
 Db 232 CTGTATAGCAGCG 291  
 QY 295 GCGTCCCTACTACAAGCAGCACTGCGCCACCGAGACCGCGCGCGCGCGCGCGCGCGCG 354  
 Db 292 GCGAGCCACTACAAGCAGCACTGCGCCCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 351  
 QY 355 ATCACCTTCGAGCGCTTCAAGGAGACCTCAAGGACTTCTCTCTCTCTCTCTCTCTCTCTCT 414

Db 352 ATCACCTTCGAGAGCTTCAAGGAGAACCTGAAGGACTTCTCTGCTGGTGATCCCTTCGAT 411  
 QY 415 TGCTGGAGCCGCTGCAGGAGT 436  
 Db 412 TGCTGGAGCCGCTGCAGGAGT 433

RESULT 7  
 AAQ97169  
 ID AAQ97169 standard; DNA; 777 BP.  
 XX  
 AC AAQ97169;  
 DT 25-AUG-1999 (first entry)  
 XX  
 DE pMON13022 DNA encoding IL-3 fusion protein.  
 XX  
 KW Interleukin-3; CSF; colony stimulating factor; cytokine; lymphokine;  
 KW mutant; mutein; fusion protein; linker; ss.  
 XX  
 OS Synthetic.  
 OS Homo sapiens.  
 XX  
 PN W09521254-A1.  
 XX  
 PD 10-AUG-1995.  
 XX  
 PF 02-FEB-1995; 95WO-US001185.  
 XX  
 PR 04-FEB-1994; 94US-00192325.  
 XX  
 PA (SEAR ) SEARLE & CO G D.  
 XX  
 PI Bauer CS, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
 PI Klein BK, Mckearn JP, Olins PO, Paik K, Thomas JW;  
 XX  
 DR WPI; 1995-283774/37.  
 DR P-PSDB; AAR79317.  
 XX  
 XX Fusion proteins comprising a human interleukin-3 variant, a linker and  
 PT interleukin-3, a variant or a colony stimulating factor - useful to  
 PT increase haematopoietic cell prodn. in a mammal.  
 XX  
 PS Claim 22; Page 158-159; 447pp; English.

A new fusion protein is disclosed which has the formula R1-L-R2, R2-L-R1,  
 R1-R2, R2-R1, R1-L-R1 or R1-R1, where R1 is a mutant or variant of human  
 interleukin-3 (hIL-3), R2 is a second colony stimulating factor (CSF)  
 including cytokine, lymphokine, interleukin, haematopoietic growth factor  
 or IL-3 variant, and L is a linker. Generic sequences are described in  
 CC AAR79235 - AAR79242, and specifically claimed examples are shown in  
 CC AAR79298-R79335 and AAR79342-R79345. The fusion protein is made by  
 CC recombinant DNA techniques. Specifically claimed examples of DNA  
 CC sequences (including the present sequence) which encode these proteins  
 CC are shown in AAQ97167-Q97204 and AAQ97222-Q97227. The fusion protein is  
 CC used to increase haematopoietic cell production. It is also useful as an  
 CC IL-3 antagonist or as a discrete antigenic fragment for production of  
 CC antibodies useful in immunoassays and immunotherapy. Antagonists are used  
 CC to block the growth of certain cancer cells and in treatment of asthma.  
 CC The fusion protein can also be used to stimulate bone marrow and blood  
 CC cell activation and growth in vitro before infusion; and to treat  
 CC diseases characterised by decreased levels of myeloid, erythroid,  
 CC lymphoid and/or megakaryocyte cells of the haematopoietic system. The  
 CC protein has the usual activity of both its component proteins, but may  
 CC have increased synergistic activity and reduced undesired side effects

Query Match 61.9%; Score 283.6; DB 2; Length 777;  
 Best Local Similarity 81.6%; Pred. No. 1.6e-42;  
 Matches 328; Conservative 0; Mismatches 74; Indels 0; Gaps 0;  
 QY 34 TCCGGCATCGAGGGCCGCAATGGCGCGCAGCGCGAGCCCGCTCCACCCAGCGGTGG 93

Db 376 TCTGGCGCGGCTCAACATGGCACCGGCTGTTCCCGTCCCGTCTACCCAGCGTGG 435  
 QY 94 GAGCAGCTGAACGCGGATCCAGGAGGCCCGCAGGCTCTCTCAACCTCTCCCGCGACACGCGC 153  
 Db 436 GAACACGTGAATGCCATCCAGGAGGCCCGCGGCTCTCTGAACCTGAGTAGAGACACTGCT 495  
 QY 154 GCCGAGATGAACGAGACCGCTGGAGGTGATCTCCGAGATGTTGATCTCCAGGAGCGGACC 213  
 Db 496 GCTGAGATGAATGAACAGTAGAAGTGATATCAGAAATTTTGGACCTCCAGGAGCGGACT 555  
 QY 214 TGCTCCAGACCGCTCGAGCTGTACAAGCAGGCGCTCGCGGCGAGCTTCACCAAGCTC 273  
 Db 556 TGCTACAGACCGCTCGAGCTGTACAAGCAGGCGCTCGCGGCGAGCTTCACCAAGCTC 615  
 QY 274 AAGGGCCCGCTCACCATGATGGCGTCCCACTACAAGCAGCAGCTGCCACCGACCGCGGAG 333  
 Db 616 AAGGGCCCGCTTGACCATGATGGCCAGCCACTACAAGCAGCAGCTGCCCTCCAACCCCGAA 675  
 QY 334 ACTCTCTGCGCCACCCAGATCATCACCCTTCGAGAGCTTCAAGGAGAACCTCAAGGACTTC 393  
 Db 676 ACTTCTGTGCAACCCAGATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTC 735  
 QY 394 CTCCTCGTGTATCCCGTTGACTGCTGGGAGCCGCTGCAGGAG 435  
 Db 736 CTGCTGTGTATCCCTTTGACTGCTGGGAGCCAGTCCAGGAG 777

RESULT 8  
 AAA03723  
 ID AAA03723 standard; DNA; 777 BP.  
 XX  
 AC AAA03723;  
 XX  
 DT 19-MAY-2000 (first entry)  
 XX  
 DE Human interleukin-3 mutant containing fusion protein DNA SEQ ID NO:55.  
 XX  
 KW Human; interleukin 3; IL-3; mutant; mutein; CSF; cytokine;  
 KW colony stimulating factor; haematopoietic growth factor; lymphokine;  
 KW fusion protein; haematopoietic disorder; infection; cancer;  
 KW radiation therapy; chemotherapy; bone marrow suppressive drug;  
 KW bone marrow activation; blood cell activation; blood transplant; ds.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 XX  
 PN US6022535-A.  
 XX  
 PD 08-FEB-2000.  
 XX  
 PF 06-JUN-1995; 95US-00469318.  
 XX  
 PR 04-FEB-1994; 94US-00192325.  
 PR 02-FEB-1995; 95WO-US001185.  
 PR 06-APR-1995; 95US-00411795.  
 XX  
 PA (SEAR ) SEARLE & CO G D.  
 XX  
 PI Bauer SC, Abrams MA, Braford-Goldberg SR, Easton AM, Klein BK;  
 PI Paik K, Thomas JW, Mckearn JP, Olins PO, Caparon MH;  
 XX  
 DR WPI; 2000-160368/14.  
 XX  
 PT Treating hematopoietic disorders with fusion proteins comprising mutated  
 PT interleukin-3 fused with secondary colony stimulating factors or other  
 PT interleukin-3 variants.  
 XX  
 PS Example 27; Col 135-136; 276pp; English.  
 XX  
 CC Methods have been developed for treating haematopoietic disorders with  
 CC fusion proteins comprising recombinant, mutated human interleukin-3 (hIL-  
 CC 3) variants or mutant proteins (muteins) fused with secondary colony

stimulating factors (CSFs) (e.g. cytokines, lymphokines, interleukin and/or haematopoietic colony stimulating factors) or other interleukin-3 variants with or without a linker. The methods may be used in vivo to treat haematopoietic disorders resulting from bacterial, viral and fungal infections, cancer radiation therapy, chemotherapy or bone marrow suppressive drugs. They may also be used in vitro to stimulate bone marrow and blood cell activation and growth prior to infusion of the bone marrow and blood transplants into patients. IL-3 is a haematopoietic growth factor which has the property of being able to promote the survival, growth and differentiation of haematopoietic cells. The fusion molecules are characterised by possessing the usual activity of both of their constituent peptides and further by having a biological or physiological activity greater than the additive function of the IL-3 or second CSF alone (i.e. the peptides act synergistically). Their activity may also be further enhanced by the mutations they comprise. The variations may further reduce undesirable side effects associated with IL-3. ANY53130 to ANY53226, and AAA03721 to AAA03782 represent sequences used in the exemplification of the present invention

Sequence 777 BP; 204 A; 227 C; 183 G; 163 T; 0 U; 0 Other;

Query Match 61.9%; Score 283.6; DB 3; Length 777;  
 Best Local Similarity 81.6%; Pred. No. 1.6e-42;  
 Matches 328; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

34 TCCGGCATCGAGGGCGCGATGCGCGCGAGCGCGCGCGCTCCACCGCGGTGG 93  
 |||||  
 376 TCTGGCGGGCTCCAAATGCGACCGCGCTCGTTCCCGCTACCCAGCGGTGG 435  
 |||||

94 GAGCAGTGAACCGGATCCAGAGGCGCGCGAGCTCTCAACTCTCCCGGACACCGCC 153  
 |||||

436 GAACAGTGAATGCCATCCAGGAGCGCGCGCTCTCTGAACCTGAGTAGACACTGCT 495  
 |||||

154 GCGAGATGAACGAGACCGTGGAGGTGATCCGAGATGTCGATCTCCAGAGCGGACC 213  
 |||||

496 GCTGAGTGAATGAACAGTAGAAGTGATCAGAAATGTTGACCTCCAGAGCGGACT 555  
 |||||

214 TGCTCTCAGACCGCGCTCGAGGTGTACAAAGCAGGGCGCTCCGGGCGAGCTCCACGAGCTC 273  
 |||||

556 TGCTACAGACCGCGCTCGAGGTGTACAAAGCAGGGCGCTCCGGGCGAGCTCCACGAGCTC 615  
 |||||

274 AGGGCGCGCTACCATGATGCGGTCCCACTACAGCAGACTGCGCCACGACCGCGAG 333  
 |||||

616 AAGGGCGCGCTTGACCATGATGCGCGAGCGCACTACAGCAGCACTGCGCTCCAAACCGCGAA 675  
 |||||

334 ACTCTGCGCGCACCCAGATCATCCTTCGAGAGCTTCAAGAGAACTCAAGGACTTC 393  
 |||||

676 ACTTCTGTGCAACCCAGATTATCCTTTGAAAGTTTCAAGAGAACTGAGGACTTC 735  
 |||||

394 CTCTCTGATCCCGTTGCACTGCTGGAGCGCGGTGACGAG 435  
 |||||

736 CTGCTGTGATCCCTTTGACTGCTGGAGCCAGTCCAGAG 777  
 |||||

## RESULT 9

ABX00012

ID ABX00012 standard; DNA; 777 BP.

XX AC ABX00012;

XX DT 18-DEC-2002 (first entry)

DE DE Human interleukin-3 associated DNA sequence #3.

XX KW Haematopoietic factor; GM-CSF; colony stimulating factor; CSF-1; ds;

XX KW G-CSF; G-CSFser17; c-mpl ligand; TPO; MGF; erythropoietin; flt3 ligand;

XX KW human growth hormone; B-cell growth factor; leukaemia;

XX KW B-cell differentiation factor; eosinophil differentiation factor;

XX KW stem cell factor; SCF; cyclic neutropenia; aplastic anaemia;

XX KW thrombocytopenia; idiopathic neutropenia; Chediak-Higashi syndrome;

XX KW systemic lupus erythematosus; SLE; myelodysplastic syndrome;

XX KW myelofibrosis; Interleukin-3; IL-3; stem cell.

OS Unidentified.

XX US6436387-B1.

PN 20-AUG-2002.

PD 09-DEC-1996; 96US-00762227.

XX 24-NOV-1992; 92US-00981044.

XX 22-NOV-1993; 93WO-US011197.

PR 04-FEB-1994; 94US-00192325.

PR 06-FEB-1995; 95WO-US001185.

PR 06-APR-1995; 95US-00411795.

PR 06-JUN-1995; 95US-00446872.

XX (SEAR ) SEARLE &amp; CO G D.

XX Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;

PI Klein BK, McKearn JP, Ollins PO, Paik K, Thomas JW;

XX WPI; 2002-749206/81.

XX Ex vivo expansion of stem cells, for enhancing transduction efficiency of

PT cultured stem cells, comprises culturing stem cells in growth medium

PT having mutant interleukin-3, and hematopoietic factor, and harvesting

PT cultured cells.

PS Disclosure; Col 161-164; 203pp; English.

XX The invention relates to ex vivo expansion of stem cells, comprises

CC culturing stem cells with a growth medium comprising a chimaera protein,

CC and harvesting the cultured stem cells. The chimaera is based on a

CC mutant human interleukin-3 (IL 3) sequence coupled to a haematopoietic

CC factor (e.g. GM-CSF (colony stimulating factor), CSF-1, G-CSF, G-

CC CSFser17, c-mpl ligand TPO, MGF, erythropoietin, IL-13, IL-15, IL-16,

CC flt3 ligand, human growth hormone, B-cell growth factor, B-cell

CC differentiation factor, eosinophil differentiation factor and stem cell

CC factor (SCP)) via a peptide linker. The formula for the chimaera is given

CC in the specification. Also included is a method for enhancing the

CC efficiency of the transduction of cultured stem cells by a heterologous

CC gene, comprising: (a) removing stem cells from a patient or donor; (b)

CC culturing the stem cells with a growth medium comprising the chimaera (c)

CC transducing DNA into cultured cells; and (d) harvesting the transduced

CC cells. The method is useful for ex vivo expansion of stem cells, and

CC enhancing the efficiency of the transduction of cultured stem cells by a

CC heterologous gene. The method is also useful for treating a patient

CC having a haematopoietic disorder. The expanded haematopoietic cells are

CC also useful in the treatment of cyclic neutropenia, aplastic anaemia,

CC thrombocytopenia, idiopathic neutropenia, Chediak-Higashi syndrome,

CC systemic lupus erythematosus (SLE), leukaemia, myelodysplastic syndrome

CC and myelofibrosis. The present sequence is an IL-3 mutant associated DNA

CC sequence. Note: The present sequence is included in the sequence listing

CC but is not mentioned anywhere else in the specification

XX

SQ Sequence 777 BP; 204 A; 227 C; 183 G; 163 T; 0 U; 0 Other;

Query Match 61.9%; Score 283.6; DB 6; Length 777;

Best Local Similarity 81.6%; Pred. No. 1.6e-42;

Matches 328; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

QY 34 TCCGGCATCGAGGGCGCGATGCGCGCGAGCGCGCGCGCTCCACCGCGGTGG 93  
 |||||

Db 376 TCTGGCGGGCTCCAAATGCGACCGCGCTCGTTCCCGCTACCCAGCGGTGG 435  
 |||||

QY 94 GAGCAGTGAACCGGATCCAGAGGCGCGCGAGCTCTCAACTCTCCCGGACACCGCC 153  
 |||||

Db 436 GAACAGTGAATGCCATCCAGGAGCGCGCGCTCTCTGAACCTGAGTAGACACTGCT 495  
 |||||

QY 154 GCGAGATGAACGAGACCGTGGAGGTGATCTCCGAGATGTCGATCTCCAGAGCGGACC 213  
 |||||

Db 496 GCTGAGTGAATGAACAGTAGAAGTGATCAGAAATGTTGACCTCCAGAGCGGACT 555  
 |||||

QY 214 TGCTCTCAGACCGCGCTCGAGCTGTACAGAGGGGCTCCGGGAGCTCCACGAGCTC 273  
 |||||

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Db      556 TGCTACAGACCCGCTGGAGCTGTACAAGCAGGGCTGGGGCAGCTCACCAAGCTC 615
Qy      274 AAGGGCCCGCTCACCATGATGGGTCCCACTAAGCAGCACTGCCACCGACCGCGAG 333
Db      616 AAGGGCCCGCTTGACATGATGGCCAGCCTACAAGCAGCACTGCCCTCCAAACCCGGAA 675
Qy      334 ACCTCTGCGCCGCCACCCAGATCATCAGCTTCGAGAGCTTCAGGAGACCTCAAGGACTTC 393
Db      676 ACTTCTGTGCAACCCAGATTATACCTTTGAAAGTTTCAAAGAGAACCTGAAGGACTTC 735
Qy      394 CTCTCTGTGATCCCGTTTCGACTGTGCGAGCCGGTGCCAGGAG 435
Db      736 CTGCTTGTATCCCTTTGACTGCTGGAGCCAGTCAGGAG 777

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## RESULT 10

ADJ14267  
ID ADJ14267 standard; DNA; 777 BP.

AC ADJ14267;

XX 20-MAY-2004 (first entry)

DT DNA related to human interleukin-3 (IL-3) mutant protein - SEQ ID 55.

DE stem cell; antianaemic; immunostimulant; immunomodulator;  
KW antiinflammatory; dermatological; immunosuppressive; cytostatic;  
KW neuroprotective; haemopoietic disorder; gene therapy; myeloid; erythroid;  
KW lymphoid; megakaryocyte; aplastic anaemia; periodic neutropenia;  
KW Chediak-Higashi syndrome; systemic lupus erythematosus; leukaemia;  
KW myelodysplastic syndrome; myelofibrosis; interleukin-3; IL-3; ds.

XX Unidentified.

XX US2003185790-A1.

XX 02-OCT-2003.

XX 26-FEB-2002; 2002US-00083446.

XX 24-NOV-1992; 92US-00981044.

PR 22-NOV-1993; 93WO-US011197.

PR 04-FEB-1994; 94US-00192325.

PR 02-FEB-1995; 95WO-US001185.

PR 06-APR-1995; 95US-00411795.

PR 06-JUN-1995; 95US-00446872.

PR 09-DEC-1996; 96US-00762227.

XX (BAUE/) BAUER S C.

PA (ABRA/) ABRAMS M A.

PA (BRA/) BRAFORD-GOLDBERG S R.

PA (CAPA/) CAPARON M H.

PA (EAST/) EASTON A M.

PA (KLEI/) KLEIN B K.

PA (MCKE/) MCKEARN J P.

PA (OLIN/) OLINS P O.

PA (PAIK/) PAIK K.

PA (THOM/) THOMAS J W.

XX Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;

PI Klein BK, Mckearn JP, Olins PO, Paik K, Thomas JW;

XX WPI; 2004-096775/10.

XX Ex vivo expansion of stem cells, e.g. hematopoietic cells for treating  
PT aplastic anemia, involves culturing the stem cells with growth medium  
PT comprising chimera protein, and harvesting the cultured stem cells.

XX Disclosure; SEQ ID NO 55; 202pp; English.

XX The invention relates to a novel method whereby stem cells are ex vivo  
CC expanded via culturing the stem cells with a growth medium comprising a

CC chimera protein, followed by harvesting of the cultured stem cells. The  
CC method of the invention has antianaemic, immunostimulant, the  
CC immunomodulator, antiinflammatory, dermatological, immunosuppressive,  
CC cytostatic and neuroprotective applications and may be useful to target  
CC haemopoietic cells for gene therapy, preferably for treating patients  
CC having a haemopoietic disorder characterised by decreased levels of  
CC myeloid, erythroid, lymphoid, and/or megakaryocyte cells of haemopoietic  
CC system. The expanded ex vivo cells may be used to treat neutropenia,  
CC aplastic anaemia, periodic neutropenia, Chediak-Higashi syndrome,  
CC systemic lupus erythematosus, leukaemia, myelodysplastic syndrome or  
CC myelofibrosis. The current sequence is that of a DNA related to the human  
CC interleukin-3 (IL-3) mutant protein of the invention.

XX Sequence 777 BP; 204 A; 227 C; 183 G; 163 T; 0 U; 0 Other;

Query Match 61.9%; Score 283.6; DB 12; Length 777;

Best Local Similarity 81.6%; Pred. NO. 1.6e-42;

Matches 328; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

Qy 34 TCCGGCATCGAGGGCCGCGATGGCGCCAGCGCGAGCCCGCGTCCACCGAGCGGTGG 93

Db 376 TCTGGCGCGGCTCCAAACATGGCACCGGCTGTTCCTCCCGTCTACCCAGCGGTGG 435

Qy 94 GAGCAGCTGAACGCGATCCAGGAGGCCCGCAGGCTCTCAACCTCTCCCGCGACACCGCC 153

Db 436 GAACAGTGAATGCCATCCAGGAGGCCCGCGTCTCTGAACCTGAGTAGACACTGCT 495

Qy 154 GCCGAGATGAACGAGACCGGTGGAGGTGATCTCCGAGATGTTTCGATCTCCAGGAGCGGACC 213

Db 496 GCTGAGATGATGAACAGTAGAAGTGATATCAGAAATGTTTACCTCCAGGAGCGGACT 555

Qy 214 TGCTCCAGACCGCTCCGAGCTGTACAAGAGGGGCTCCGCGGAGGCTTCACAAGCTC 273

Db 556 TGCCTACAGACCGCGCTGGAGCTGTACAAGCAGGGCTTCGCGGCGAGCTTCACCAAGCTC 615

Qy 274 AAGGGCCCGCTCACCATGATGGCGTCCCACTACAAGCAGCACTGCCACCGACCGCGAG 333

Db 616 AAGGGCCCGCTTGACCATGATGGCCAGCCACTACAAGCAGCACTGCCCTCCAAACCGCGAA 675

Qy 334 ACCTCTGCGCCACCCAGATCATCAGCTTCGAGAGCTTCAAGGAGAACCTCAAGGACTTC 393

Db 676 ACTTCTGTGCAACCCAGATTATCAGCTTTGAAAGTTTCAAAGAGAACCTGAAGGACTTC 735

Qy 394 CTCCTCGTATCCCGTTTCGACTGCTGGAGCCGGTGCCAGGAG 435

Db 736 CTGCTTGTATCCCTTTGACTGCTGGAGCCAGTCAGGAG 777

## RESULT 11

AAQ97208

ID AAQ97208 standard; DNA; 402 BP.

AC AAQ97208;

XX 25-AUG-1999 (first entry)

DT pMONI3012 DNA sequence.

XX Interleukin; hIL-3; CSF; colony stimulating factor; cytokine; lymphokine;

XX mutant; mutein; fusion protein; linker; ss.

XX Synthetic.

OS Homo sapiens.

XX WO9521254-A1.

XX 10-AUG-1995.

XX 02-FEB-1995; 95WO-US001185.

XX 04-FEB-1994; 94US-00192325.

XX (SEAR ) SEARLE & CO G D.





QY 172 GTGGAGTGTATCCAGATGTTGATCTCCAGGAGCCGACCTGCTCCAGACCCGCTC 231  
 DB 121 GTAGAAGTGTATCAGAAATGTTGACCTCCAGGAGCCGACTTGCCTACAGACCCGCTG 180  
 QY 232 GAGCTGTACAAGCAGGCGCTCCGCGCAGCTCACCAGCTCAAGGCGCGCTCACCATG 291  
 DB 181 GAGCTGTACAAGCAGGCGCTCCGCGCAGCTCACCAGCTCAAGGCGCGCTCACCATG 240  
 QY 292 ATGGCGTCCCACTACAAGCAGCACTCCCAAGCAGCCGAGACCTCTCCGCGCACCCAG 351  
 DB 241 ATGGCCAGCCACTACAAGCAGCACTCCCAAGCAGCCGAGACCTCTCCGCGCACCCAG 300  
 QY 352 ATCATCACCCTTGAGAGCTTCAAGGAGAACCTCAAGGACTTCTCTCGTATCCGCTTC 411  
 DB 301 ATTATCACCTTTGAAAGTTTCAAGAGAAACCTGAAGGACTTCTCTGCTATCCCTTT 360  
 QY 412 GACTGCTGGGAGCGGTGCAGGAGTCA 438  
 DB 361 GACTGCTGGGAGCGGTGCAGGAGTGA 387

## RESULT 13

ABX00086  
 ID ABX00086 standard; DNA; 402 BP.

AC ABX00086;

DT 18-DEC-2002 (first entry)

DE Human interleukin-3 associated DNA sequence #77.

OS Haematopoietic factor; GM-CSF; colony stimulating factor; CSF-1; ds;  
 KW G-CSF; G-CSFser17; c-mpl ligand; TPO; MGDF; erythropoietin; flt3 ligand;  
 KW human growth hormone; B-cell growth factor; leukaemia;  
 KW B-cell differentiation factor; eosinophil differentiation factor;  
 KW stem cell factor; SCF; cyclic neutropenia; aplastic anaemia;  
 KW thrombocytopenia; idiopathic neutropenia; Chediak-Higashi syndrome;  
 KW systemic lupus erythematosus; SLE; myelodysplastic syndrome;  
 KW myelofibrosis; Interleukin-3; IL-3; stem cell.

XX Unidentified.

XX US6436387-B1.

PN 20-AUG-2002.

XX 09-DEC-1996; 96US-00762227.

XX 24-NOV-1992; 92US-00981044.

PR 22-NOV-1993; 93WO-US011197.

PR 04-FEB-1994; 94US-00192325.

PR 04-FEB-1995; 95WO-US001185.

PR 06-APR-1995; 95US-00411795.

XX 06-JUN-1995; 95US-00446872.

XX (SEAR ) SEARLE & CO G D.

XX Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;

PI Klein BK, Mckearn JP, Oline PO, Paik K, Thomas JW;

XX WPI; 2002-749206/81.

XX Ex vivo expansion of stem cells, for enhancing transduction efficiency of

PT cultured stem cells, comprises culturing stem cells in growth medium

PT having mutant interleukin-3, and hematopoietic factor, and harvesting

PT cultured cells.

XX Disclosure; Col 299-300; 203pp; English.

XX The invention relates to ex vivo expansion of stem cells, comprises

CC mutated human interleukin-3 (IL-3) sequence coupled to a haematopoietic  
 CC factor (e.g. GM-CSF (colony stimulating factor), CSF-1, G-CSF, G-  
 CC CSFser17, c-mpl ligand TPO, MGDF, erythropoietin, IL-13, IL-15, IL-16,  
 CC flt3 ligand, human growth hormone, B-cell growth factor, B-cell  
 CC differentiation factor, eosinophil differentiation factor and stem cell  
 CC factor (SCF)) via a peptide linker. The formula for the chimera is given  
 CC in the specification. Also included is a method for enhancing the  
 CC efficiency of the transduction of cultured stem cells by a heterologous  
 CC gene, comprising: (a) removing stem cells from a patient or donor; (b)  
 CC culturing the stem cells with a growth medium comprising the chimera (c)  
 CC transducing DNA into cultured cells; and (d) harvesting the transduced  
 CC cells. The method is useful for ex vivo expansion of stem cells, and  
 CC enhancing the efficiency of the transduction of cultured stem cells by a  
 CC heterologous gene. The method is also useful for treating a patient  
 CC having a haematopoietic disorder. The expanded haematopoietic cells are  
 CC also useful in the treatment of cyclic neutropenia, aplastic anaemia,  
 CC thrombocytopenia, idiopathic neutropenia, Chediak-Higashi syndrome,  
 CC systemic lupus erythematosus (SLE), leukaemia, myelodysplastic syndrome  
 CC and myelofibrosis. The present sequence is an IL-3 mutant associated DNA  
 CC sequence. Note: The present sequence is included in the sequence listing  
 CC but is not mentioned anywhere else in the specification  
 XX  
 SQ Sequence 402 BP; 99 A; 124 C; 101 G; 78 T; 0 U; 0 Other;

Query Match 61.8%; Score 283; DB 6; Length 402;

Best Local Similarity 83.2%; Pred. No. 2.1e-42;

Matches 322; Conservative 0; Mismatches 65; Indels 0; Gaps 0;

QY 52 ATGGCGCCAGCGCGAGCCGCTCCACCCAGCCGCTGGAGCAGCTGAACGGGATC 111

DB 1 ATGGCACCGGCTGCTTCCCGTCCCGTACCCAGCGCTGGAGCAGCTGAATGCCATC 60

QY 112 CAGGAGGCGCGAGGCTCTCTCAACCTCTCCCGCAGCACCGCCGCGAGATGAACGAGCC 171

DB 61 CAGGAGGCGCGGCTCTCTCAACCTCTAGTAGAGACACTGCTGCTGAGATGAATGAACA 120

QY 172 GTGGAGGTGATCTCGAGATGTTGATCTCCAGGAGCGACCTGCTCCAGACCCGCTC 231

DB 121 GTAGAAGTGTATCAGAAATGTTTGAACCTCCAGGAGCGACTTGGCTACAGACCCGCTG 180

QY 232 GAGCTGTACAAGCAGGCGCTCCGCGCAGCTCACCAGCTCAAGGGCGCTCACCATG 291

DB 181 GAGCTGTACAAGCAGGCGCTCCGCGCAGCTCACCAGCTCAAGGGCGCTTGCACATG 240

QY 292 ATGGCGTCCCACTACAAGCAGCACTGCCACCGACCCGCGAGACCTCTCCGCCACCCAG 351

DB 241 ATGGCCAGCCACTACAAGCAGCACTGCCCTCAACCCCGGAAACTTCTCTGTGCAACCCAG 300

QY 352 ATCATCACCCTTCGAGAGCTTCAAGGAGAACCTCAAGGACTTCTCTCTGATCCCGTTC 411

DB 301 ATTATCACCTTTGAAAGTTTCAAGAGAAACCTGAAGGACTTCTCTGCTATCCCTTT 360

QY 412 GACTGCTGGGAGCGGTGCAGGAGTGA 438

DB 361 GACTGCTGGGAGCGGTGCAGGAGTGA 387

## RESULT 14

ADJ14388

ID ADJ14388 standard; DNA; 402 BP.

XX ADJ14388;

XX ADJ14388;

DT 20-MAY-2004 (first entry)

XX DNA related to human interleukin-3 (IL-3) mutant protein - SEQ ID 151.

XX stem cell; antianaemic; immunostimulant; immunomodulator;

KW antiinflammatory; dermatological; immunosuppressive; cytostatic;

KW neuroprotective; haematopoietic disorder; gene therapy; myeloid; erythroid;

KW lymphoid; megakaryocyte; aplastic anaemia; periodic neutropenia;

KW Chediak-Higashi syndrome; systemic lupus erythematosus; leukaemia;

KW myelodysplastic syndrome; myelofibrosis; interleukin-3; IL-3; ds.

XX OS Unidentified.  
XX PN US2003185790-A1.  
XX PD 02-OCT-2003.  
XX PF 26-FEB-2002; 2002US-00083446.  
XX PR 24-NOV-1992; 92US-00981044.  
XX PR 22-NOV-1993; 93WO-US011197.  
XX PR 04-FEB-1994; 94US-00192325.  
XX PR 02-FEB-1995; 95WO-US0001185.  
XX PR 06-APR-1995; 95US-00411795.  
XX PR 06-JUN-1995; 95US-00446872.  
XX PR 09-DEC-1996; 96US-00762227.  
XX PA (BAUE/) BAUER S C.  
XX PA (ABRA/) ABRAMS M A.  
XX PA (BRAB/) BRAFORD-GOLDBERG S R.  
XX PA (CAPA/) CAPARON M H.  
XX PA (EAST/) EASTON A M.  
XX PA (KLEI/) KLEIN B K.  
XX PA (MCKE/) MCKEARN J P.  
XX PA (OLIN/) OLINS P O.  
XX PA (PAIK/) PAIK K.  
XX PA (THOM/) THOMAS J W.  
XX PI Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM,  
XX PI Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;  
XX XX WPI; 2004-096775/10.  
XX DR Ex vivo expansion of stem cells, e.g. hematopoietic cells for treating  
XX PT aplastic anemia, involves culturing the stem cells with growth medium  
XX PT comprising chimera protein, and harvesting the cultured stem cells.  
XX XX Disclosure; SEQ ID NO 176; 202pp; English.  
XX CC The invention relates to a novel method whereby stem cells are ex vivo  
XX CC expanded via culturing the stem cells with a growth medium comprising a  
XX CC chimera protein, followed by harvesting of the cultured stem cells. The  
XX CC method of the invention has antianemic, immunostimulant,  
XX CC immunomodulator, antiinflammatory, dermatological, immunosuppressive,  
XX CC cytostatic and neuroprotective applications and may be useful to target  
XX CC hematopoietic cells for gene therapy, preferably for treating patients  
XX CC having a haemopoietic disorder characterised by decreased levels of  
XX CC myeloid, erythroid, lymphoid, and/or megakaryocyte cells of haemopoietic  
XX CC system. The expanded ex vivo cells may be used to treat neutropenia,  
XX CC aplastic anaemia, periodic neutropenia, Chediak-Higashi syndrome,  
XX CC systemic lupus erythematosus, leukaemia, myelodysplastic syndrome or  
XX CC myelofibrosis. The current sequence is that of a DNA related to the human  
XX CC interleukin-3 (IL-3) mutant protein of the invention.  
XX SQ Sequence 402 BP; 99 A; 124 C; 101 G; 78 T; 0 U; 0 Other;  
Query Match 61.8%; Score 283; DB 12; Length 402;  
Best Local Similarity 83.2%; Pred. No. 2.1e-42;  
Matches 322; Conservative 0; Mismatches 65; Indels 0; Gaps 0;  
QY 52 ATGGCCGACGCGGAGCGGAGCCCGTCCACCCAGCCGTGGAGACGTCGACCGGATC 111  
DB 1 ATGGCACCAGGCTGTTCCCGCTCCCGCTTACCAGCCGTGGGAAACACGTGAATGCATC 60  
QY 112 CAGGAGCCGCGAGGCTCTCAACTCTCCCGGACACCGCCGAGATGAACGAGCC 171  
DB 61 CAGGAGCCGCGGCGTCTCTTGAACTGTAGTAGACACTGCTGCTGAGATGAATGAACA 120  
QY 172 GTGGAGGTGATCTCCGAGATGTTTCGATCTCCAGGAGCCGACCTGCTCCAGACCCCGCTTC 231  
DB 121 GTAGAAGTGATATCAGAAATGTTTGACCTCCAGGAGCCGACTGCTCAGACCCCGCTG 180  
QY 232 GAGCTGTACAGCAGGCGCTCCGCGGACGCTCCACCAAGCTCAAGGGCCGCTCACCATG 291

DB 181 GAGCTGTACAGCAGGCGCTCGGGGAGCCCTCAACCAAGCTCAAGGGCCCTTGACCATG 240  
QY 292 ATGGCGTCCCACTACAAGCAGCACTGCCACCGACCCCGGAGACCTCTCTGCCGCCACCCAG 351  
DB 241 ATGGCCAGCCACTACAAGCAGCACTGCCCTCAACCCCGGAAACTTCTGTGCAACCCAG 300  
QY 352 ATCATCACCTTCGAGAGCTTCAAGGAGAACTCAAGGACTTCTCTCTCGTATCCGCTTC 411  
DB 301 ATTATCACCTTTGAAAGTTTCAAGAGAACTTCAAGGAGACTTCTCTGTGTCATCCCTTT 360  
QY 412 GACTCTGGGAGCCGCTGCAGGAGTGA 438  
DB 361 GACTCTGGGAGCCAGTCCAGGAGTGA 387  
RESULT 15  
AAQ97183  
ID AAQ97183 standard; DNA; 822 BP.  
XX AC AAQ97183;  
XX DT 25-AUG-1999 (first entry)  
XX DE PMON13035 DNA encoding IL-3 fusion protein.  
XX KW Interleukin; hIL-3; CSF; colony stimulating factor; cytokine; lymphokine;  
XX KW mutant; mutein; fusion protein; linker; ss.  
XX OS Synthetic.  
XX OS Homo sapiens.  
XX PN WO9521254-A1.  
XX PD 10-AUG-1995.  
XX PF 02-FEB-1995; 95WO-US0001185.  
XX PR 04-FEB-1994; 94US-00192325.  
XX (SEAR ) SEARLE & CO G D.  
XX PI Bauer CS, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
XX PI Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;  
XX XX WPI; 1995-283774/37.  
XX P-PSDB; AAR79320.  
XX PT Fusion proteins comprising a human interleukin-3 variant, a linker and  
XX PT interleukin-3, a variant or a colony stimulating factor - useful to  
XX PT increase haematopoietic cell prodn. in a mammal.  
XX PS Claim 22; Page 167-168; 447pp; English.  
XX CC A new fusion protein is disclosed which has the formula R1-L-R2, R2-L-R1,  
XX CC R1-R2, R2-R1, R1-L-R1 or R1-R1, where R1 is a mutant or variant of human  
XX CC interleukin-3 (hIL-3), R2 is a second colony stimulating factor (CSF)  
XX CC including cytokine, lymphokine, interleukin, haematopoietic growth factor  
XX CC or IL-3 variant, and L is a linker. Generic sequences are described in  
XX CC AAR03225 AAR03242, and specifically claimed examples are shown in  
XX CC AAR79298-R79335 and AAR79342-R79345. The fusion protein is made by  
XX CC recombinant DNA techniques. Specifically claimed examples of DNA  
XX CC sequences (including the present sequence) which encode these proteins  
XX CC are shown in AAQ97167-Q97204 and AAQ97222-Q97227. The fusion protein is  
XX CC used to increase haematopoietic cell production. It is also useful as an  
XX CC IL-3 antagonist or as a discrete antigenic fragment for production of  
XX CC antibodies useful in immunoassays and immunotherapy. Antagonists are used  
XX CC to block the growth of certain cancer cells and in treatment of asthma.  
XX CC The fusion protein can also be used to stimulate bone marrow and blood  
XX CC cell activation and growth in vitro before infusion; and to treat  
XX CC diseases characterised by decreased levels of myeloid, erythroid,  
XX CC lymphoid and/or megakaryocyte cells of the haematopoietic system. The  
XX CC protein has the usual activity of both its component proteins, but may



CC have increased synergistic activity and reduced undesired side effects

XX Sequence 822 BP; 222 A; 244 C; 180 G; 176 T; 0 U; 0 Other;

Query Match 61.4%; Score 281; DB 2; Length 822;  
Best Local Similarity 83.1%; Pred. No. 4.8e-42;  
Matches 320; Conservative 0; Mismatches 65; Indels 0; Gaps 0;

Qy	51	CATGGCGCCAGCGCGAGCCGCGGAGCCGCTCCACCCAGCCGCTGGAGCAGCTGAACGGAT	110
Db	438	CATGGCACCGGCTCGTTCCCGCTCCCGCTACCCAGCCGCTGGGAACACGCTGAATGCCAT	497
Qy	111	CCAGGAGGCCCGCAGGCTCCTCAACCTCTCCCGCGACACCGCGCGGAGATGAACGAGAC	170
Db	498	CCAGGAGGCCCGCGCTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAATGAAC	557
Qy	171	CGTGGAGGTGATCTCCGAGATGTTGATCTCCAGGAGCCGACCTGCTCCAGACCCGCT	230
Db	558	AGTAGAAGTGATATCAGAAATGTTGACCTCCAGGAGCCGACTTGCTACAGACCCGCT	617
Qy	231	CGAGCTGTACAGCAGGGCTCCGCGGAGCCTCAGCAAGCTCAAGGGCCGCTACCAT	290
Db	618	GGAGCTGTACAGCAGGGCTCCGCGGAGCCTCAGCAAGCTCAAGGGCCGCTTGACCAT	677
Qy	291	GATGGCGTCCCACTACAAGCAGCACTGCCACCGCGGAGACCTCTCTGGCCACCCA	350
Db	678	GATGGCCAGCCACTACAAGCAGCACTGCCCTCAACCCCGGAACTTCCTGTGCAACCCA	737
Qy	351	GATCATCACCTTCGAGAGCTTCAAGGAGAACCTCAAGGACTTCCTCTCTGATCCGTT	410
Db	738	GATTATCACCTTTGAAAGTTTCAAGAGAACCTGAGGACTTCCTGCTGTATCCCTT	797
Qy	411	CGACTGCTGGGAGCCGCTGAGGAG	435
Db	798	TGACTGCTGGGAGCCAGTCCAGGAG	822

Search completed: March 11, 2005, 15:58:59  
Job time : 425 secs



GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: March 11, 2005, 16:42:11 ; Search time 453 Seconds  
(without alignments)  
6014.183 Million cell updates/sec

Title: us-10-723-083-1

Perfect score: 458

Sequence: 1 cggccggatgcaccacca.....gtcgtgcgcatgcgcg 458

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 5537552 seqs, 2974263231 residues

Total number of hits satisfying chosen parameters: 11075104

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:

- 1: /cgn2\_6/ptodata/1/pubpna/US07\_PUBCOMB.seq:
- 2: /cgn2\_6/ptodata/1/pubpna/PCT\_NEW\_PUB.seq:
- 3: /cgn2\_6/ptodata/1/pubpna/US06\_NEW\_PUB.seq:
- 4: /cgn2\_6/ptodata/1/pubpna/US06\_PUBCOMB.seq:
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- 7: /cgn2\_6/ptodata/1/pubpna/US08\_NEW\_PUB.seq:
- 8: /cgn2\_6/ptodata/1/pubpna/US08\_PUBCOMB.seq:
- 9: /cgn2\_6/ptodata/1/pubpna/US09A\_PUBCOMB.seq:
- 10: /cgn2\_6/ptodata/1/pubpna/US09B\_PUBCOMB.seq:
- 11: /cgn2\_6/ptodata/1/pubpna/US09C\_PUBCOMB.seq:
- 12: /cgn2\_6/ptodata/1/pubpna/US09\_NEW\_PUB.seq:
- 13: /cgn2\_6/ptodata/1/pubpna/US10A\_PUBCOMB.seq:
- 14: /cgn2\_6/ptodata/1/pubpna/US10B\_PUBCOMB.seq:
- 15: /cgn2\_6/ptodata/1/pubpna/US10C\_PUBCOMB.seq:
- 16: /cgn2\_6/ptodata/1/pubpna/US10D\_PUBCOMB.seq:
- 17: /cgn2\_6/ptodata/1/pubpna/US10E\_PUBCOMB.seq:
- 18: /cgn2\_6/ptodata/1/pubpna/US10F\_PUBCOMB.seq:
- 19: /cgn2\_6/ptodata/1/pubpna/US10\_NEW\_PUB.seq:
- 20: /cgn2\_6/ptodata/1/pubpna/US11\_NEW\_PUB.seq:
- 21: /cgn2\_6/ptodata/1/pubpna/US60\_NEW\_PUB.seq:
- 22: /cgn2\_6/ptodata/1/pubpna/US60\_PUBCOMB.seq:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	458	100.0	458	19	US-10-723-083-1
2	318.6	69.6	429	19	US-10-723-083-3
3	312.6	68.3	435	17	US-10-188-056-31
4	311	67.9	435	17	US-10-188-056-33
5	283.6	61.9	777	16	US-10-083-446-55
6	283	61.8	402	16	US-10-083-446-176
7	281	61.4	822	16	US-10-083-446-69
8	281	61.4	903	16	US-10-083-446-66
9	277	60.5	2211	17	US-10-609-346-9
10	276.8	60.4	448	17	US-10-609-346-19
11	276.8	60.4	781	17	US-10-447-315-20
					Sequence 1, Appl
					Sequence 3, Appl
					Sequence 31, Appl
					Sequence 33, Appl
					Sequence 55, Appl
					Sequence 176, Appl
					Sequence 69, Appl
					Sequence 66, Appl
					Sequence 9, Appl
					Sequence 19, Appl
					Sequence 20, Appl

12	276.8	60.4	789	16	US-10-131-985-16	Sequence 16, Appl
13	276.8	60.4	789	19	US-10-901-417-16	Sequence 16, Appl
14	276.2	60.3	1011	13	US-10-044-090-509	Sequence 509, Appl
15	275.8	60.2	435	9	US-09-826-025-8	Sequence 8, Appl
16	275.8	60.2	435	14	US-10-083-590-14	Sequence 14, Appl
17	275.8	60.2	435	17	US-10-411-037-17	Sequence 17, Appl
18	275.8	60.2	435	17	US-10-411-026-17	Sequence 17, Appl
19	275.8	60.2	435	17	US-10-410-962-17	Sequence 17, Appl
20	275.8	60.2	435	17	US-10-411-049-17	Sequence 17, Appl
21	275.8	60.2	435	18	US-10-410-930-17	Sequence 17, Appl
22	275.8	60.2	435	18	US-10-410-997-17	Sequence 17, Appl
23	275.8	60.2	435	18	US-10-411-012-17	Sequence 17, Appl
24	275.8	60.2	435	18	US-10-287-994-17	Sequence 17, Appl
25	275.8	60.2	435	18	US-10-410-913-17	Sequence 17, Appl
26	275.8	60.2	435	18	US-10-785-577-8	Sequence 8, Appl
27	275.8	60.2	435	19	US-10-410-980-17	Sequence 17, Appl
28	275.2	60.1	505	18	US-10-688-845-82	Sequence 82, Appl
29	275.2	60.1	737	15	US-10-081-969-19	Sequence 19, Appl
30	275.2	60.1	756	15	US-10-177-390-21	Sequence 21, Appl
31	275.2	60.1	756	17	US-10-351-157-180	Sequence 180, App
32	275.2	60.1	756	17	US-10-352-554-165	Sequence 165, App
33	275.2	60.1	756	17	US-10-429-802-32	Sequence 32, Appl
34	275.2	60.1	756	17	US-10-430-503-23	Sequence 23, Appl
35	275.2	60.1	756	17	US-10-305-720-1195	Sequence 1195, Ap
36	275.2	60.1	756	18	US-10-475-024-7	Sequence 7, Appl
37	275.2	60.1	756	18	US-10-475-024-8	Sequence 8, Appl
38	275.2	60.1	767	18	US-10-666-122-4	Sequence 4, Appl
39	275.2	60.1	767	18	US-10-666-122-6	Sequence 6, Appl
40	275.2	60.1	767	19	US-10-278-698-30	Sequence 30, Appl
41	275.2	60.1	767	19	US-10-278-698-30	Sequence 30, Appl
42	275.2	60.1	1318	14	US-10-228-811-3	Sequence 544, App
43	274.8	60.0	579	17	US-10-449-831A-187	Sequence 3, Appl
44	274.2	59.9	435	17	US-10-188-056-32	Sequence 187, App
45	274.2	59.9	496	16	US-10-267-384-191	Sequence 191, Appl

ALIGNMENTS

RESULT 1

US-10-723-083-1  
; Sequence 1, Application US/10723083  
; Publication No. US2005050602A1  
; GENERAL INFORMATION:  
; APPLICANT: Altosar, Illimar  
; APPLICANT: Sardana, Ravinder  
; APPLICANT: Dudani, Aail  
; APPLICANT: Ganz, Peter  
; APPLICANT: Tackaberry, Elleen  
; TITLE OF INVENTION: Production of GM-CSF in Plants  
; FILE REFERENCE: 08-898901US  
; CURRENT APPLICATION NUMBER: US/10/723,083  
; CURRENT FILING DATE: 2003-11-26  
; PRIOR APPLICATION NUMBER: Canada 2,410,702  
; PRIOR FILING DATE: 2002-11-26  
; NUMBER OF SEQ ID NOS: 4  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 1  
; LENGTH: 458  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: (10)..(438)  
; OTHER INFORMATION:  
US-10-723-083-1

Query Match 100.0%; Score 458; DB 19; Length 458;

Best Local Similarity 100.0%; Pred. No. 7.2e-110;

Matches 458; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

1 CGGCCGGATGCACCACACACACTCTCCGGCATCGAGGGCCGCGCCCA 60

Db 1 CGGCCGGGATGCACCAACACACACCACTCTCTCCGGGATCGAGGGCCGATGGCGCA 60  
QY 61 GCGCGAGGCCGAGCCGCTCCACCCAGCCGCTGGAGACAGTGAAACCGGATCCAGGAGGCC 120  
Db 61 GCGCGAGGCCGAGCCGCTCCACCCAGCCGCTGGAGACAGTGAAACCGGATCCAGGAGGCC 120  
QY 121 CGCAGGCTCTCAACCTCTCCCGGACACCGCCGCGGAGATGAACGAGACCGTGGAGGTG 180  
Db 121 CGCAGGCTCTCAACCTCTCCCGGACACCGCCGCGGAGATGAACGAGACCGTGGAGGTG 180  
QY 181 ATCTCCGAGATGTTGATCTCCAGGAGCCGACTGCTCCAGACCCGCTCCAGCTGTAC 240  
Db 181 ATCTCCGAGATGTTGATCTCCAGGAGCCGACTGCTCCAGACCCGCTCCAGCTGTAC 240  
QY 241 AAGCAGGCTCCGCGGACCTCACCAGCTCAAGGCGCCGCTCACCATGATGGCGTCC 300  
Db 241 AAGCAGGCTCCGCGGACCTCACCAGCTCAAGGCGCCGCTCACCATGATGGCGTCC 300  
QY 301 CACTACAGCAGACTGCCACCGACCGCCGAGACCTCTCGGCCACCCAGATCATCAC 360  
Db 301 CACTACAGCAGACTGCCACCGACCGCCGAGACCTCTCGGCCACCCAGATCATCAC 360  
QY 361 TTCGAGAGCTTCAAGGAGACCTCAAGGACTTCTCTCTGATCCGTTGACTGCTGG 420  
Db 361 TTCGAGAGCTTCAAGGAGACCTCAAGGACTTCTCTCTGATCCGTTGACTGCTGG 420  
QY 421 GAGCCGGTGAGGAGTGAAGTACGCTAGCGTACGATGCGG 458  
Db 421 GAGCCGGTGAGGAGTGAAGTACGCTAGCGTACGATGCGG 458

## RESULT 2

US-10-723-083-3  
; Sequence 3, Application US/10723083  
; Publication No. US2005005062A1  
; GENERAL INFORMATION:  
; APPLICANT: Alcosaar, Illimar  
; APPLICANT: Sardan, Ravinder  
; APPLICANT: Dudani, Aail  
; APPLICANT: Ganz, Peter  
; APPLICANT: Tackberry, Eileen  
; TITLE OF INVENTION: Production of GM-CSF in Plants  
; FILE REFERENCE: 08-98901US  
; CURRENT APPLICATION NUMBER: US/10/723,083  
; CURRENT FILING DATE: 2003-11-26  
; PRIOR APPLICATION NUMBER: Canada 2,410,702  
; PRIOR FILING DATE: 2002-11-26  
; NUMBER OF SEQ ID NOS: 4  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 3  
; LENGTH: 429  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: (1)..(429)  
; OTHER INFORMATION:  
US-10-723-083-3

Query Match 59.6%; Score 318.6; DB 19; Length 429;  
Best Local Similarity 83.9%; Pred. No. 1.6e-73;  
Matches 360; Conservative 0; Mismatches 69; Indels 0; Gaps 0;

QY 10 ATGCACACACACACACCACTCTCTCCGGGATCGAGGGCCGATGGCGCGCAGC 69  
Db 1 ATGCACACACACACACCACTCTCTCCGGGATCGAGGGCCGATGGCGCGCAGC 60  
QY 70 CGAGGCCGCTCAACCGAGCGTGGAGACAGTGAAACCGATCAGAGGCCGCGAGGCTC 129  
Db 61 CCAGGCCGAGCAGCGAGCGCTGGAGCATGTGAATGCCATCAGAGGCCGCGGCTCTC 120  
QY 130 CTCACCTCTCCGCGACACCGCCGCGAGATGAACGAGACCGTGGAGGTATCCAG 189

Db 121 CTGAACCTTGAGTAGAGACACTCTCTCTGAGATGAATGAACAGTAGAGTGAATCAGAA 180  
QY 190 ATGTTTCGATCTCCAGGAGCCGACCTGCTCCAGACCCGCTCGAGCTGTACAAGCAGGCG 249  
Db 181 ATGTTTCGATCTCCAGGAGCCGACCTGCTCCAGACCCGCTCGAGCTGTACAAGCAGGCG 240  
QY 250 CTCGCGGAGGCTCACCAGCTCAAGGGCCGCTCAGCATGATGGCTCCACCTACAG 309  
Db 241 CTCGCGGAGGCTCACCAGCTCAAGGGCCGCTTGGACCATGATGGCCAGCCACTACAG 300  
QY 310 CAGCACTGCCACCCGAGACCTCTCTGCGGACCTCTGCGCCACCCAGATCATCACCTTCAGAGC 369  
Db 301 CAGCACTGCCCTCCAAACCCGGAATCTCTCTGTGCAACCCAGATTAACCTTTGAAAGT 360  
QY 370 TTCAAGAGAACTCAAGGACTTCTCTCTGATCCGTTGAGTGTGGGAGCGGCTG 429  
Db 361 TTCAAGAGAACTCAAGGACTTCTCTCTGATCCCTTGTCTATCCCTTGTGCTGGGAGCCAGTC 420  
QY 430 CAGGAGTGA 438  
Db 421 CAGGAGTGA 429

## RESULT 3

US-10-188-056-31  
; Sequence 31, Application US/10188056  
; Publication No. US20040009934A1  
; GENERAL INFORMATION:  
; APPLICANT: Qiu, Jian-Tai  
; APPLICANT: Lai, Wan-Ching  
; APPLICANT: Chu, Yong Liang  
; APPLICANT: Li, Frank Q.  
; TITLE OF INVENTION: Improved GM-CSF Nucleic Acid Sequences  
; FILE REFERENCE: 3781-004-27  
; CURRENT APPLICATION NUMBER: US/10/188,056  
; CURRENT FILING DATE: 2002-09-26  
; NUMBER OF SEQ ID NOS: 34  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 31  
; LENGTH: 435  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-188-056-31

Query Match 68.3%; Score 312.6; DB 17; Length 435;  
Best Local Similarity 82.8%; Pred. No. 5.7e-72;  
Matches 357; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

QY 8 GGATGCACACACACACCACTCTCTCGGATCGAGGGCCGATGGCGCGCAGCGCGA 67  
Db 5 GGCTGCAGAGCTGTCTCTGTGGGCACCGTGGCTGCAGCATCAGCGCTCCCGCCAGAA 64  
QY 68 GCGCGAGCCGCTCCACCCAGCGCTGGGAGCAGCTGAACCGATCCAGAGGCCCGCAGCG 127  
Db 65 GCGCGAGCCGCTCCACCCAGCGCTGGGAGCAGCTGAACCGATCCAGAGGCCCGCAGCG 124  
QY 128 TCCTCAACCTCTCCCGGACACCGCGCGGAGATGAACGAGACCGTGGAGGTGATCTCG 187  
Db 125 TGCTGAACCTGTCCAGAGACACCGCGCGAGATGAACGAGACCGTGGAGGTGATCAGG 184  
QY 188 AGATTTTGATCTCAGGAGCGGAGCTGCTCTCCAGACCCGCTCGAGCTGTACAAGCAGG 247  
Db 185 AGATTTTGATCTCAGGAGCGGAGCTGCTCTCCAGACCCGCTGGAGCTGTACAAGCAGG 244  
QY 248 GCCTCCGCGGAGCTCACCAGCTCAAGGGCCGCTCACCATGATGGCTCCCACTACA 307  
Db 245 GACTCGGGGAGCTGACCAAGCTGAAGGAGCCCTGACCATGATGGCCAGCCACTACA 304  
QY 308 AGCAGCACTGCCCAACCGGAGGAGCTCTGTGGCCACCCAGATCATCACTTTCGAGA 367  
Db 305 AGCAGCACTGCCCTCCACACCCGAGACAGCTGTGGCCACCCAGATCATCACTTTCGAGA 364  
QY 368 GCTTCAAGAGNACCTCAAGGACTTCTCTCTGATCCCGTTGATCCCGTGGGAGCGG 427

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Db      365 GCTTCAGGAGACCTGAAGGACTTCTCTGCTGGTGATCCCTTCGACTGCTGGAGCCG 424
Qy      428 TGCAGGAGTGA 438
Db      425 TGCAGGAGTGA 435

RESULT 4
US-10-188-056-33
; Sequence 33, Application US/10188056
; Publication No. US2004009934A1
; GENERAL INFORMATION:
; APPLICANT: Qiu, Jian-Tai
; APPLICANT: Lai, Wan-Ching
; APPLICANT: Chu, Yong Liang
; APPLICANT: Li, Frank Q.
; TITLE OF INVENTION: Improved GM-CSF Nucleic Acid Sequences
; FILE REFERENCE: 3781-004-27
; CURRENT APPLICATION NUMBER: US/10/188,056
; CURRENT FILING DATE: 2002-09-26
; NUMBER OF SEQ ID NOS: 34
; SOFTWARE: Fast-Seq for Windows Version 4.0
; SEQ ID NO 33
; LENGTH: 435
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-188-056-33

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Query Match      67.9%; Score 311; DB 17; Length 435;
Best Local Similarity 82.6%; Pred. No. 1.5e-71;
Matches 356; Conservative 0; Mismatches 75; Indels 0; Gaps 0;

Qy      8 GGATGCACCAACCAACCACTCTCTCCGGCATCGAGGCGCGCATGCGCCAGCGCGCA 67
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Qy      68 GCCGAGCGCGTCAACCAAGCGTGGAGAGCACTGTGAACGCGATCCAGAGGCGCGCAGGC 127
Db      65 GCCCGAGCGCGTCAACCAAGCGTGGAGAGCACTGTGAACGCGATCCAGAGGCGCGCAGGC 124

Qy      128 TCCTCAACCTCTCCGGGACACCGCGCGAGATGACGAGCGGTGGAGTGATCTCG 187
Db      125 TGTGAACCTGTTCAGAGACACCGCGCGAGATGACGAGCGGTGGAGTGATCAGCG 184

Qy      188 AGATGTTGAGTCTCCAGAGCGCGACCTGTCTCCAGACCGCGCTCGAGCTGTACAAGCAGG 247
Db      185 AGATGTTGAGTCTCCAGAGCGCGACCTGTCTCCAGACCGCGCTCGAGCTGTACAAGCAGG 244

Qy      248 GCCTCCGCGGAGCTTCAACCAAGCTCAAGGCGCGCTCAACCATGATGGGTCCCACTACA 307
Db      245 GACTGGGGGAGCGCTGACCAAGCTGAAGGAGCGCGCTGACCATGATGGCGCAGCACTACA 304

Qy      308 AGCAGACTCTCCACCGCGCGAGACTCTCTCCGCGCGAGATCATCATCATCATCATCAT 367
Db      305 AGCAGACTCTCCACCGCGCGAGACTCTCTCCGCGCGAGATCATCATCATCATCATCAT 364

Qy      368 GCTTCAAGGAGAACTTCAAGGAGTCTCTCTCTGATGATGATGATGATGATGATGATG 427
Db      365 GCTTCAAGGAGAACTTCAAGGAGTCTCTCTCTGATGATGATGATGATGATGATGATG 424

Qy      428 TGCAGGAGTGA 438
Db      425 TGCAGGAGTGA 435

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RESULT 5
US-10-083-446-55
; Sequence 55, Application US/10083446
; Publication No. US20030185790A1
; GENERAL INFORMATION:
; APPLICANT: Abrams, Mark A.
;           Bauer, S. C.

```

```

; Braford-Goldberg, Sarah R.
; Caparon, Mairé H.
; Easton, Alan M.
; Klein, Barbara K.
; McKearn, John P.
; Olin, Peter O.
; Paik, Kumnan
; Thomas, John W.

TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells
Using Multivariant (IL-3) Hematopoiesis Chimera Proteins
NUMBER OF SEQUENCES: 197
CORRESPONDENCE ADDRESS:
ADDRESSES: S. Christopher Bauer, Pharmacia Corporation
STREET: 800 N. Lindbergh
CITY: St. Louis
STATE: Missouri
COUNTRY: USA
ZIP: 63167

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.25
CURRENT APPLICATION DATA: US/10/083,446
APPLICATION NUMBER: US/10/083,446
FILING DATE: 26-Feb-2002
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/762,227
FILING DATE: 09-DEC-1996
APPLICATION NUMBER: US 08/192,325
FILING DATE: 14-FEB-1994
APPLICATION NUMBER: US 08/446,872
FILING DATE: 06-JUN-1995
ATTORNEY/AGENT INFORMATION:
NAME: S. Christopher Bauer
REGISTRATION NUMBER: 42,305
REFERENCE/DOCKET NUMBER: C-2790/6
TELECOMMUNICATION INFORMATION:
TELEPHONE: (636)737-6257
TELEFAX: (636)737-5452
INFORMATION FOR SEQ ID NO: 55:
SEQUENCE CHARACTERISTICS:
LENGTH: 777 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
SEQUENCE DESCRIPTION: SEQ ID NO: 55:
US-10-083-446-55

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Query Match      61.9%; Score 283.6; DB 16; Length 777;
Best Local Similarity 81.6%; Pred. No. 2e-64;
Matches 328; Conservative 0; Mismatches 74; Indels 0; Gaps 0;

Qy      34 TCGGAGTCGAGGCGCGCATGCGCGCGAGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 93
Db      376 TCTGGCGGGGCTTCAACATGCGCGCGCTCGTTCCCGCTCCCGCTACCGAGCGGTGG 435

Qy      94 GAGCAGTGAACGCGCATGCGAGGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 153
Db      436 GAACAGTGAATGCCATCCAGGAGGCGCGCGCGCTCTCTGAACTGATGATGATGATGAT 495

Qy      154 GCGGAGTGAACGAGACCGTGGAGTGATCTCCGATGTTGATGTTGATGTTGATGTTGATG 213
Db      496 GCTGATGAATGAACAGTAGAAGTATCAGAAATGTTGATGTTGATGTTGATGTTGATG 555

Qy      214 TGCCTCCAGACCGCGCTCGAGTGTACAAGAGGCGCGCGCGCGCGCGCGCGCGCGCGCG 273
Db      556 TGCCTACAGACCGCGCTCGAGTGTACAAGAGGCGCGCGCGCGCGCGCGCGCGCGCGCG 615

Qy      274 AAGGGCCCGCTCACCATGATGCGGTCCCTACTACAAGAGCATGTGCCACCGACCGCGGAG 333

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Db 616 AAGGGCCCTTGACCATGATGGCCAGCCACTACAAGCAGCACTGCCTCCAAACCCGGAA 675  
QY 334 ACCTCTGCGCCACCCAGATCATCACTTCGAGAGCTTCAAGGAGAACTCAAGGACTTC 393  
Db 676 ACTTCTGTGCAACCCAGATTAICACTTTGAAAGTTTCAAGAGAACTTGAAGGACTTC 735  
QY 394 CTCTCGTATCCCGTTTCGACTCTCTGGAGCCGGTGCAGAG 435  
Db 736 CTGCTTGTATCCCTTTGACTCTCTGGAGCCAGTCCAGAG 777

RESULT 6  
US-10-083-446-176  
; Sequence 176, Application US/10083446  
; Publication No. US20030185790A1  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; Bauer, S. C.  
; Braford-Goldberg, Sarah R.  
; Caparon, Mairé H.  
; Easton, Alan M.  
; Klein, Barbara K.  
; McKearn, John P.  
; Ollins, Peter O.  
; Paik, Kuman  
; Thomas, John W.  
; TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells  
; Using Multivariant (IL-3) Hematopoiesis Chimera Proteins  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: S. Christopher Bauer, Pharmacia Corporation  
; Corporate Patent Dept., Mail Zone O4E  
; STREET: 800 N. Lindbergh  
; CITY: St. Louis  
; STATE: Missouri  
; COUNTRY: USA  
; ZIP: 63167  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/10/083,446  
; FILING DATE: 26-Feb-2002  
; CLASSIFICATION: <Unknown>  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/762,227  
; FILING DATE: 09-DEC-1996  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; APPLICATION NUMBER: US 08/446,872  
; FILING DATE: 06-JUN-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: S. Christopher Bauer  
; REGISTRATION NUMBER: 42,305  
; REFERENCE/DOCKET NUMBER: C-2790/6  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (636)737-6257  
; TELEFAX: (636)737-5452  
; INFORMATION FOR SEQ ID NO: 176:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 402 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; SEQUENCE DESCRIPTION: SEQ ID NO: 176:  
US-10-083-446-176  
Query Match 61.8%; Score 283; DB 16; Length 402;  
Best Local Similarity 83.2%; Pred. No. 3e-64;

Matches 322; Conservative 0; Mismatches 65; Indels 0; Gaps 0;  
QY 52 ATGGCGCCAGCCGCGAGCCCGAGCCCGTCCACCCAGCCGTGGAGCAGCTGAACGCGATC 111  
Db 1 ATGGCACCAGCTCGTTCCCGTCCCGTCTACCCAGCCGTGGGAACACGTAATGCCATC 60  
QY 112 CAGGAGGCCGCGAGCGCTCTCTCAACCTCTCCCGCGACAACCGCCGCGAGATGAACGAGACC 171  
Db 61 CAGGAGGCCGCGCGCTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAATGAACA 120  
QY 172 GTGGAGGTGATCTCCGAGATGTTCCATCTCCAGGAGCCGACCTGCTCCAGACCCGCTC 231  
Db 121 GTAGAAAGTATATCAGAAATGTTGACCTCCAGGAGCCGACTTGCCTACAGACCCGCTG 180  
QY 232 GAGCTGTACAAGCAGGGCGCTCCCGCGAGCCCTCACAAGCTCAAGGGCCCGCTCACCATG 291  
Db 181 GAGCTGTACAAGCAGGGCGCTCCCGCGAGCCCTCACAAGCTCAAGGGCCCGCTTGCATG 240  
QY 292 ATGGCGTCCACTACAAGCAGCACTGCGCCACCGACCCCGGAGACCTCTGCGCCACCCAG 351  
Db 241 ATGGCCAGCCACTACAAGCAGCACTGCGCTCCCAACCCCGGAAACTTCTGTGCAACCCAG 300  
QY 352 ATCATCAGCTTCGAGAGCTTCAAGGAGAACCTCAAGGACTTCTCTCGTGATCCGCTTC 411  
Db 301 ATTATCAGCTTTGAAAGTTTCAAGAGAACTTCAAGAGAACTTCCCTGCTTGTATCCCTTT 360  
QY 412 GACTGCTGGAGCCCGTGCAGGAGTGA 438  
Db 361 GACTGCTGGAGCCAGTCCAGGAGTGA 387

RESULT 7  
US-10-083-446-69  
; Sequence 69, Application US/10083446  
; Publication No. US20030185790A1  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; Bauer, S. C.  
; Braford-Goldberg, Sarah R.  
; Caparon, Mairé H.  
; Easton, Alan M.  
; Klein, Barbara K.  
; McKearn, John P.  
; Ollins, Peter O.  
; Paik, Kuman  
; Thomas, John W.  
; TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells  
; Using Multivariant (IL-3) Hematopoiesis Chimera Proteins  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: S. Christopher Bauer, Pharmacia Corporation  
; Corporate Patent Dept., Mail Zone O4E  
; STREET: 800 N. Lindbergh  
; CITY: St. Louis  
; STATE: Missouri  
; COUNTRY: USA  
; ZIP: 63167  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/10/083,446  
; FILING DATE: 26-Feb-2002  
; CLASSIFICATION: <Unknown>  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/762,227  
; FILING DATE: 09-DEC-1996  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; APPLICATION NUMBER: US 08/446,872  
; FILING DATE: 06-JUN-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: S. Christopher Bauer  
; REGISTRATION NUMBER: 42,305  
; REFERENCE/DOCKET NUMBER: C-2790/6  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (636)737-6257  
; TELEFAX: (636)737-5452  
; INFORMATION FOR SEQ ID NO: 176:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 402 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; SEQUENCE DESCRIPTION: SEQ ID NO: 176:  
US-10-083-446-176  
Query Match 61.8%; Score 283; DB 16; Length 402;  
Best Local Similarity 83.2%; Pred. No. 3e-64;



; GENERAL INFORMATION:  
; APPLICANT: Yu, Zailin  
; APPLICANT: Fu, Yan  
; TITLE OF INVENTION: RECOMBINANT HUMAN ALBUMIN FUSION PROTEINS WITH LONG-LASTING BIOLOGICAL ACTIVITY  
; TITLE OF INVENTION: EFFECTS  
; FILE REFERENCE: ZYU-0603  
; CURRENT APPLICATION NUMBER: US/10/609,346  
; CURRENT FILING DATE: 2003-06-26  
; PRIOR APPLICATION NUMBER: US 60/392,948  
; PRIOR FILING DATE: 2002-07-01  
; NUMBER OF SEQ ID NOS: 40  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 9  
; LENGTH: 2211  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: DNA of HSA-GMCSF  
US-10-609-346-9

Query Match 60.5%; Score 277; DB 17; Length 2211;  
Best Local Similarity 82.0%; Pred. No. 1.1e-62;  
Matches 319; Conservative 0; Mismatches 70; Indels 0; Gaps 0;

QY 50 GCATGGGGCCAGCGCGGAGCGGCGGCTCCACCGAGCGGTGGAGCAGCTGAACGGGA 109  
DB 1823 GCTTAGCACCGCGCGCTCGCCAGCGCCAGCAGCGCCCTGGAGCATGTGAATGCCA 1882  
QY 110 TCCAGGAGGGCCGACGGCTCCTCAACTCTCCCGGACACCGCGCGGAGATGAACGAGA 169  
DB 1883 TCCAGGAGGGCCGCGGCTCCTCAACTCTGAGTAGAGACACTGCTGTGATGAATGAAA 1942  
QY 170 CCGTGGAGGTGATCTCCGAGATGTTGATCTCCAGGAGCGGAGCTGCTCTCCAGACCGCGC 229  
DB 1943 CAGTAGAGTCACTCAGAAATGTTTGACCTCCAGGAGCGGAGCTGCTCTACAGACCGCGC 2002  
QY 230 TCGAGCTGTGAACGACGAGGGCTCCGGGCGAGCTCAACAGCTCAAGGGCGCGCTCAACA 289  
DB 2003 TGGAGCTGTGAACGACGAGGGCTCCGGGCGAGCTCAACAGCTCAAGGGCGCGCTCAACA 2062  
QY 290 TGAATGGGTCCCACTACAGCAGCACTGCCCCACGAGCCCGGAGACCTCTGCGCCACCC 349  
DB 2063 TGAATGGGTCCCACTACAGCAGCACTGCCCCACGAGCCCGGAGACCTCTGCGCCACCC 2122  
QY 350 AGATCATCACTTCGAGAGCTTCAAGAGAGAACTCAAGGACTTCTCTCTGTCATCCCGT 409  
DB 2123 AGATTCATCACTTTGAAAGTTTCAAGAGAACTCAAGGACTTCTCTCTGTCATCCCGT 2182  
QY 410 TCGACTGCTGGAGCGCGGTCCAGAGTGA 438  
DB 2183 TTGACTGCTGGAGCGCGGTCCAGAGTGA 2211

RESULT 10  
US-10-609-346-19  
; Sequence 19, Application US/10609346  
; Publication No. US20040063635A1  
; GENERAL INFORMATION:  
; APPLICANT: Yu, Zailin  
; APPLICANT: Fu, Yan  
; TITLE OF INVENTION: RECOMBINANT HUMAN ALBUMIN FUSION PROTEINS WITH LONG-LASTING BIOLOGICAL ACTIVITY  
; TITLE OF INVENTION: EFFECTS  
; FILE REFERENCE: ZYU-0603  
; CURRENT APPLICATION NUMBER: US/10/609,346  
; CURRENT FILING DATE: 2003-06-26  
; PRIOR APPLICATION NUMBER: US 60/392,948  
; PRIOR FILING DATE: 2002-07-01  
; NUMBER OF SEQ ID NOS: 40  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 19  
; LENGTH: 448  
; TYPE: DNA  
; ORGANISM: Homo sapiens



QY 128 TCCTCAACCTCTCCCGGACACCGCCGCGAGATGAACGAGACCGTGGAGGTGATCTCCG 187  
DB 157 TCCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAATGAACAGTAGAAGTCATCTCAG 216  
QY 188 AGATGTTTCGATCTCCAGGAGCCGACCTGCTCCAGACCCGCTCGAGCTGTACAAGCAGG 247  
DB 217 AAATGTTTGAACCTCCAGGAGCCGACCTGCTCAAGACCCGCTGGAGCTGTACAAGCAGG 276  
QY 248 GCCTCCGCGGACCTCACCAGCTCAAGGCGCCGCTCACCATGATGGGCTCCCACTACA 307  
DB 277 GCCTGCGGGGACGCTCACCAGCTCAAGGCGCCCTTGACCATGATGGCCAGCCACTACA 336  
QY 308 AGCAGACTGCCCCACCGACCCCGGAGACCTCTCTGCGCCACCCAGATCATCACTTCGAGA 367  
DB 337 AGCAGACTGCCCCACCGACCCCGGAACTTCTGTGTCAACCCAGATTATCACTTTGAAA 396  
QY 368 GCTTCAAGGAGACCTCAAGGACTTCTCTCTGTGATCCGTTTCGACTGCTGGAGCCGG 427  
DB 397 GTTTCAAAGAGAACCTGAAGGACTTCTGCTGTGATCCCTTTGACTGCTGGAGCCAG 456  
QY 428 TGCAGGAGTGAG 439  
DB 457 TCCAGGAGTGAG 468

## RESULT 12

US-10-131-985-16  
; Sequence 16, Application US/10131985  
; Publication No. US20030199440A1  
; GENERAL INFORMATION:  
; APPLICANT: Dack, Kevin N  
; APPLICANT: Davies, Michael J  
; APPLICANT: Fish, Paul V  
; APPLICANT: Huggins, Jonathan P  
; APPLICANT: McIntosh, Nicholas L  
; APPLICANT: Occleston, Nicholas L  
; TITLE OF INVENTION: Composition  
; FILE REFERENCE: PCS 10391A  
; CURRENT APPLICATION NUMBER: US/10/131,985  
; CURRENT FILING DATE: 2002-04-25  
; PRIOR APPLICATION NUMBER: US/09/726,295  
; PRIOR FILING DATE: 2000-11-30  
; PRIOR APPLICATION NUMBER: GB 9930768.8  
; PRIOR FILING DATE: 1999-12-29  
; NUMBER OF SEQ ID NOS: 60  
; SOFTWARE: PatentIn Ver. 2.1  
; SEQ ID NO 16  
; LENGTH: 789  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-131-985-16

Query Match 60.4%; Score 276.8; DB 16; Length 789;  
Best Local Similarity 77.5%; Pred. No. 1.2e-62;  
Matches 335; Conservative 0; Mismatches 97; Indels 0; Gaps 0;  
QY 8 GGATGCACCAACCAACCACTCTCTCGGCATCGAGGCGCGCATGGCGCCAGCGCGCA 67  
DB 37 GGCTGAGAGCCTGTGCTCTTGGGCACTGTGGCCCTGCGAGCATCTCTGACCCGCCGCT 96  
QY 68 GCCGAGCCCGTCCACCCAGCCGTCGGAGCACGCTGAACCGCATCCAGGAGGCCCGCAGGC 127  
DB 97 CGCCAGCCCCAGCAGCGCCCTGGAGCATGTGAATGCCATCCAGGAGGCCCGCGTC 156  
QY 128 TCCTCAACCTCTCCCGGACACCGCCGCGAGATGAACGAGACCGTGGAGGTGATCTCCG 187  
DB 157 TCCTGAACCTGAGTAGAGACACTGCTGTGATGAATGAATGAACAGTAGAAGTCATCTCAG 216  
QY 188 AGATGTTTCGATCTCCAGGAGCCGACCTGCTCCAGACCCGCTCGAGCTGTACAAGCAGG 247  
DB 217 AAATGTTTGAACCTCCAGGAGCCGACCTGCTACAGACCCGCTGGAGCTGTACAAGCAGG 276  
QY 248 GCCTCCGCGGACCTCAACAGCTCAAGGCGCCGCTCAACATGATGGCGTCCCACTACA 307  
DB 157 TCCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAATGAACAGTAGAAGTCATCTCAG 216  
QY 188 AGATGTTTCGATCTCCAGGAGCCGACCTGCTCCAGACCCGCTCGAGCTGTACAAGCAGG 247  
DB 217 AAATGTTTGAACCTCCAGGAGCCGACCTGCTACAGACCCGCTGGAGCTGTACAAGCAGG 276

QY 248 GCCTCCGCGGACCTCACCAGCTCAAGGCGCCGCTCACCATGATGGGTCCCACTACA 307  
DB 277 GCCTGCGGGGACGCTCACCAGCTCAAGGCGCCCTTGACCATGATGGCCAGCCACTACA 336  
QY 308 AGCAGACTGCCCCACCGACCCCGGAGACCTCTCTGCGCCACCCAGATCATCACTTCGAGA 367  
DB 337 AGCAGACTGCCCCACCGACCCCGGAACTTCTGTGTCAACCCAGATTATCACTTTGAAA 396  
QY 368 GCTTCAAGGAGACCTCAAGGACTTCTCTCTGTGATCCGTTTCGACTGCTGGAGCCGG 427  
DB 397 GTTTCAAAGAGAACCTGAAGGACTTCTGCTGTGATCCCTTTGACTGCTGGAGCCAG 456  
QY 428 TGCAGGAGTGAG 439  
DB 457 TCCAGGAGTGAG 468

## RESULT 13

US-10-901-417-16  
; Sequence 16, Application US/10901417  
; Publication No. US20050026836A1  
; GENERAL INFORMATION:  
; APPLICANT: Dack, Kevin N  
; APPLICANT: Davies, Michael J  
; APPLICANT: Fish, Paul V  
; APPLICANT: Huggins, Jonathan P  
; APPLICANT: McIntosh, Fraser S  
; APPLICANT: Occleston, Nicholas L  
; TITLE OF INVENTION: Composition  
; FILE REFERENCE: PCS 10391A  
; CURRENT APPLICATION NUMBER: US/10/901,417  
; CURRENT FILING DATE: 2004-07-28  
; PRIOR APPLICATION NUMBER: US/10/131,985  
; PRIOR FILING DATE: 2002-04-25  
; PRIOR APPLICATION NUMBER: US/09/726,295  
; PRIOR FILING DATE: 2000-11-30  
; PRIOR APPLICATION NUMBER: GB 9930768.8  
; PRIOR FILING DATE: 1999-12-29  
; NUMBER OF SEQ ID NOS: 60  
; SOFTWARE: PatentIn Ver. 2.1  
; SEQ ID NO 16  
; LENGTH: 789  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-901-417-16

Query Match 60.4%; Score 276.8; DB 19; Length 789;  
Best Local Similarity 77.5%; Pred. No. 1.2e-62;  
Matches 335; Conservative 0; Mismatches 97; Indels 0; Gaps 0;  
QY 8 GGATGCACCAACCAACCACTCTCTCGGCATCGAGGCGCGCATGGCGCCAGCGCGCA 67  
DB 37 GGCTGAGAGCCTGTGCTCTTGGGCACTGTGGCCCTGCGAGCATCTCTGACCCGCCGCT 96  
QY 68 GCCGAGCCCGTCCACCCAGCCGTCGGAGCACGCTGAACCGCATCCAGGAGGCCCGCAGGC 127  
DB 97 CGCCAGCCCCAGCAGCGCCCTGGAGCATGTGAATGCCATCCAGGAGGCCCGCGTC 156  
QY 128 TCCTCAACCTCTCCCGGACACCGCCGCGAGATGAACGAGACCGTGGAGGTGATCTCCG 187  
DB 157 TCCTGAACCTGAGTAGAGACACTGCTGTGATGAATGAATGAACAGTAGAAGTCATCTCAG 216  
QY 188 AGATGTTTCGATCTCCAGGAGCCGACCTGCTCCAGACCCGCTCGAGCTGTACAAGCAGG 247  
DB 217 AAATGTTTGAACCTCCAGGAGCCGACCTGCTACAGACCCGCTGGAGCTGTACAAGCAGG 276  
QY 248 GCCTCCGCGGACCTCAACAGCTCAAGGCGCCGCTCAACATGATGGCGTCCCACTACA 307  
DB 277 GCCTGCGGGGACGCTCAACAGCTCAAGGCGCCCTTGACCATGATGGCCAGCCACTACA 336  
QY 308 AGCAGACTGCCCCACCGACCCCGGAGACCTCTCTGCGCCACCCAGATCATCACTTCGAGA 367  
DB 337 AGCAGACTGCCCCCAACCCCGGAACTTCTGTGTGCAACCCAGATTATCACTTTGAAA 396



Search completed: March 11, 2005, 18:28:48  
Job time : 457 secs

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GenCore version 5.1.6  
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OM protein - nucleic search, using frame\_plus\_p2n model

Run on: March 11, 2005, 17:24:26 ; Search time 473 Seconds  
(without alignments)  
1777.174 Million cell updates/sec

Title: US-10-723-083-2

Perfect score: 765

Sequence: 1 MHHHHSSGIEGRMAPARS.....ENLKDFLLVPPDCWEPVQE 142

Scoring table:

BLOSUM62  
Xgapop 10.0 , Xgapext 0.5  
Ygapop 10.0 , Ygapext 0.5  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

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-O=/cpn2.1/USPTO.spool/US10723083/runat\_08032005.131715.10414/app.query.fasta\_1.327  
-DB=N\_Geneseq 16Dec04 -QFMT=fastp -SUFFIX=ring -MINMATCH=0.1 -LOOPEL=0  
-LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi  
-LIST=45 -DOCALIGN=200 -THR\_SCORE=pct -THR\_MAX=100 -THR\_MIN=0 -ALIGN=15  
-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000  
-USER=US10723083 @CGN 1 1 644 @runat\_08032005.131715.10414 -NCPU=6 -ICPU=3  
-NO MAP -LARGEQUERY -NEG\_SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG  
-DSV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6  
-FGAPEXT=7 -XGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : N\_Geneseq 16Dec04:\*  
1: Geneseqn1980s:\*  
2: Geneseqn1990s:\*  
3: Geneseqn2000s:\*  
4: Geneseqn2001as:\*  
5: Geneseqn2001bs:\*  
6: Geneseqn2002as:\*  
7: Geneseqn2002bs:\*  
8: Geneseqn2003as:\*  
9: Geneseqn2003bs:\*  
10: Geneseqn2003cs:\*  
11: Geneseqn2003ds:\*  
12: Geneseqn2004as:\*  
13: Geneseqn2004bs:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	682	89.2	777	2	AaQ97169 pMON13022
2	682	89.2	777	3	Aaa03723 Human int
3	682	89.2	777	6	Abx00012 Human int
4	682	89.2	777	12	AdJ14267 DNA relat
5	679	88.8	1610	2	Aat72724 p53-GM-CS

#### ALIGNMENTS

RESULT 1  
AAQ97169  
ID AAQ97169 standard; DNA; 777 BP.  
XX  
AC AAQ97169;  
XX  
DT 25-AUG-1999 (first entry)  
XX  
DE pMON13022 DNA encoding IL-3 fusion protein.  
XX  
KW Interleukin; hIL-3; CSF; colony stimulating factor; cytokine; lymphokine;  
KW mutant; mutein; fusion protein; linker; ss.  
XX  
XX Synthetic.  
OS Homo sapiens.  
XX  
FN WO9521254-A1.  
XX  
PD 10-AUG-1995.  
XX  
PF 02-FEB-1995; 95WO-US001185.  
XX  
PR 04-FEB-1994; 94US-00192325.  
XX  
XX (SEAR ) SEARLE & CO G D.  
XX  
XX AaQ97169 pMON13022  
XX  
PI Bauer CS, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
PI Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;  
XX

DR WPI: 1995-283774/37.  
 XX P-PSDB; AAR79317.  
 XX Fusion proteins comprising a human interleukin-3 variant, a linker and  
 PT interleukin-3, a variant or a colony stimulating factor - useful to  
 PT increase haematopoietic cell prodn. in a mammal.  
 PS  
 XX Claim 22; Page 158-159; 447pp; English.  
 XX A new fusion protein is disclosed which has the formula R1-L-R2, R2-L-R1,  
 CC R1-R2, R2-R1, R1-L-R1 or R1-R1, where R1 is a mutant or variant of human  
 CC interleukin-3 (hIL-3), R2 is a second colony stimulating factor (CSF)  
 CC including cytokine, lymphokine, interleukin, haematopoietic growth factor  
 CC or IL-3 variant, and L is a linker. Generic sequences are described in  
 CC AA032335 - AA032342, and specifically claimed examples are shown in  
 CC AAR79298-R79335 and AAR79342-R79345. The fusion protein is made by  
 CC recombinant DNA techniques. Specifically claimed examples of DNA  
 CC sequences (including the present sequence) which encode these proteins  
 CC are shown in AA097167-Q97204 and AA097222-Q97227. The fusion protein is  
 CC used to increase haematopoietic cell production. It is also useful as an  
 CC IL-3 antagonist or as a discrete antigenic fragment for production of  
 CC antibodies useful in immunoassays and immunotherapy. Antagonists are used  
 CC to block the growth of certain cancer cells and in treatment of asthma.  
 CC The fusion protein can also be used to stimulate bone marrow and blood  
 CC cell activation and growth in vitro before infusion; and to treat  
 CC diseases characterised by decreased levels of myeloid, erythroid,  
 CC lymphoid and/or megakaryocyte cells of the haematopoietic system. The  
 CC protein has the usual activity of both its component proteins, but may  
 CC have increased synergistic activity and reduced undesired side effects  
 XX  
 SQ Sequence 777 BP; 204 A; 227 C; 183 G; 163 T; 0 U; 0 Other;

## Alignment Scores:

Pred. No.: 1.38e-67 Length: 777  
 Score: 682.00 Matches: 130  
 Percent Similarity: 97.01% Conservative: 0  
 Best Local Similarity: 97.01% Mismatches: 4  
 Query Match: 89.15% Indels: 0  
 DB: 2 Gaps: 0

US-10-723-083-2 (1-142) x AAQ97169 (1-777)

QY 9 SerGlylleuGluArgMetAlaProAlaArgSerProSerProSerThrGlnProTrp 28  
 DB 376 TCTGGCGGGCGCTCCAACTGGACGGCGCTCGTTCCTCCCGTCTACCCAGCGGTGG 435  
 QY 29 GluHisValAsnAlaIleGlnGluAlaArgAtcLeuLeuAsnLeuSerArgAspThrAla 48  
 DB 436 GAACACGTGATGATCCATCCAGGAGGCGCGCGTCTCTGAACCTGAGTAGACACTGTCT 495  
 QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnGluProThr 68  
 DB 496 GCTGAGATGAATCAACACAGTAGAAGTGATATCAGAAATGTTGACCTCCAGGAGCGCAT 555  
 QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88  
 DB 556 TGCTCTACAGACCGCGCTGGAGCTGTGTCAAGCAGGCGCTCGCGGCGAGCTCCACCAAGCTC 615  
 QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProProThrProGlu 108  
 DB 616 AAGGGCCCTTGACATGATGGCCAGCCACTACAGCAGACTGCGCTCCACCCCGGAA 675  
 QY 109 ThrSerCysAlaThrGlnIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128  
 DB 676 ACTTCCTGTGCAACCCAGATTATCCTTTGAAAGTTTCAAGAGAACCTGAGGACTTC 735  
 QY 129 LeuLeuValIleProPheAspCysTrpGluProValGlnGlu 142  
 DB 736 CTGCTGTTCATCCCTTTGACTGCTCGGAGCCAGTCCAGGAG 777

RESULT 2

AAA03723

ID AAA03723 standard; DNA; 777 BP.

XX AAA03723;  
 AC 19-MAY-2000 (first entry)  
 DT Human interleukin-3 mutant containing fusion protein DNA SEQ ID NO:55.  
 DE  
 XX  
 XX Human; interleukin 3; IL-3; mutant; mutein; CSF; cytokine;  
 KW colony stimulating factor; haematopoietic growth factor; lymphokine;  
 KW fusion protein; haematopoietic disorder; infection; cancer;  
 KW radiation therapy; chemotherapy; bone marrow suppressive drug;  
 XX bone marrow activation; blood cell activation; blood transplant; ds.  
 OS Homo sapiens.  
 OS Synthetic.  
 XX US6022535-A.  
 PN 08-FEB-2000.  
 XX 06-JUN-1995; 95US-00469318.  
 XX 04-FEB-1994; 94US-00192325.  
 PR 02-FEB-1995; 95WO-US001185.  
 PR 06-APR-1995; 95US-00411795.  
 XX (SEAR ) SEARLE & CO G D.  
 XX Bauer SC, Abrams MA, Brafard-Goldberg SR, Easton AM, Klein BK;  
 PI Paik K, Thomas JW, McKearn JP, Olins PO, Caparon MH;  
 XX WPI; 2000-160368/14.  
 DR Treating hematopoietic disorders with fusion proteins comprising mutated  
 XX interleukin-3 fused with secondary colony stimulating factors or other  
 XX interleukin-3 variants.  
 XX Example 27; Col 135-136; 276pp; English.  
 XX Methods have been developed for treating haematopoietic disorders with  
 CC fusion proteins comprising recombinant, mutated human interleukin-3 (hIL-  
 CC 3) variants or mutant proteins (muteins) fused with secondary colony  
 CC stimulating factors (CSFs) (e.g. cytokines, lymphokines, interleukin  
 CC and/or haematopoietic colony stimulating factors) or other interleukin-3  
 CC variants with or without a linker. The methods may be used in vivo to  
 CC treat haematopoietic disorders resulting from bacterial, viral and fungal  
 CC infections, cancer radiation therapy, chemotherapy or bone marrow  
 CC suppressive drugs. They may also be used in vitro to stimulate bone  
 CC marrow and blood cell activation and growth prior to infusion of the bone  
 CC marrow and blood transplants into patients. IL-3 is a haematopoietic  
 CC growth factor which has the property of being able to promote the  
 CC survival, growth and differentiation of haematopoietic cells. The fusion  
 CC molecules are characterised by possessing the usual activity of both of  
 CC their constituent peptides and further by having a biological or  
 CC physiological activity greater than the additive function of the IL-3 or  
 CC second CSF alone (i.e. the peptides act synergistically). Their activity  
 CC may also be further enhanced by the mutations they comprise. The  
 CC variations may further reduce undesirable side effects associated with IL  
 CC -3. AA053130 to AA053226, and AA03721 to AA03782 represent sequences  
 CC used in the exemplification of the present invention  
 XX  
 SQ Sequence 777 BP; 204 A; 227 C; 183 G; 163 T; 0 U; 0 Other;

## Alignment Scores:

Pred. No.: 1.38e-67 Length: 777  
 Score: 682.00 Matches: 130  
 Percent Similarity: 97.01% Conservative: 0  
 Best Local Similarity: 97.01% Mismatches: 4  
 Query Match: 89.15% Indels: 0  
 DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x AAA03723 (1-777)

QY 9 SerGlyLeuGluGlyArgMetAlaProAlaArgSerProSerThrGlnProTyr 28  
 Db 376 TCTGGCGCGCGCTCCAACTGGACCGCTCGTTCCCGCTCCCGCTACCCAGCGGTGG 435  
 QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAla 48  
 Db 436 GAACAGTGAATGCCATCCAGAGAGGCCCGCGCTCTCTGAACCTGAGTAGAGACACTGCT 495  
 QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnGluProThr 68  
 Db 496 GCTGAGATGAATGAACAGTAGAAGTGAATCAAGAAATGTTTGCACCTCCAGAGCGCACT 555  
 QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88  
 Db 556 TGCCTACAGACCGCTGGAGCTGTACAGAGCGGCTCGGGGCGAGCCTCACCAAGCTC 615  
 QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProThrProGlu 108  
 Db 616 AAGGGCCCTTGACCATGATGGCCAGCCACTACAGCAGCACTGCCCTCCAAACCCCGAA 675  
 QY 109 ThrSerCysAlaThrGlnIleThrPheGluSerPheLysGluLeuLysAspPhe 128  
 Db 676 ACTTCTGTGCAACCCAGATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTC 735  
 QY 129 LeuLeuValIleProPheAspCysTyrGluProValGlnGlu 142  
 Db 736 CTGCTTGTATCCCTTTGACTGCTGGAGCCAGTCCAGGAG 777

# RESULT 3 ABX00012

ID ABX00012 standard; DNA; 777 BP.

XX AC ABX00012;  
 XX 18-DEC-2002 (first entry)  
 XX Human interleukin-3 associated DNA sequence #3.  
 XX Haematopoietic factor; GM-CSF; colony stimulating factor; CSF-1; ds;  
 KW G-CSF; G-CSFser17; c-mpl ligand; TPO; MGDF; erythropoietin; flt3 ligand;  
 KW human growth hormone; B-cell growth factor; leukaemia;  
 KW B-cell differentiation factor; eosinophil differentiation factor;  
 KW stem cell factor; SCF; cyclic neutropenia; aplastic anaemia;  
 KW thrombocytopenia; idiopathic neutropenia; Chediak-Higashi syndrome;  
 KW systemic lupus erythematosus; SLE; myelodysplastic syndrome;  
 KW myelofibrosis; Interleukin-3; IL-3; stem cell.  
 XX Unidentified.  
 XX US6436387-B1.  
 XX 20-AUG-2002.  
 XX 09-DEC-1996; 96US-00762227.  
 XX 24-NOV-1992; 92US-00981044.  
 XX 22-NOV-1993; 93WO-US011197.  
 XX 04-FEB-1994; 94US-00192325.  
 XX 04-FEB-1995; 95WO-US0001185.  
 XX 06-APR-1995; 95US-00411795.  
 XX 06-JUN-1995; 95US-00446872.  
 XX (SEAR ) SEARLE & CO G D.

XX Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
 PI Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;  
 XX WPI; 2002-749206/81.

XX Ex vivo expansion of stem cells, for enhancing transduction efficiency of  
 PT cultured stem cells, comprises culturing stem cells in growth-medium  
 PT having mutant interleukin-3, and hematopoietic factor, and harvesting  
 PT cultured cells.

XX Disclosure; Col 161-164; 203pp; English.

XX The invention relates to ex vivo expansion of stem cells, comprises  
 CC culturing stem cells with a growth medium comprising a chimera protein,  
 CC and harvesting the cultured stem cells. The chimera is based on a  
 CC mutated human interleukin-3 (IL-3) sequence coupled to a haematopoietic  
 CC factor (e.g. GM-CSF (colony stimulating factor), CSF-1, G-CSF, G-  
 CC CSFser17, c-mpl ligand TPO, MGDF, erythropoietin, IL-1-13, IL-15, IL-16,  
 CC flt3 ligand, human growth hormone, B-cell growth factor, B-cell  
 CC differentiation factor, eosinophil differentiation factor and stem cell  
 CC factor (SCF)) via a peptide linker. The formula for the chimera is given  
 CC in the specification. Also included is a method for enhancing the  
 CC efficiency of the transduction of cultured stem cells by a heterologous  
 CC gene, comprising: (a) removing stem cells from a patient or donor; (b)  
 CC culturing the stem cells with a growth medium comprising the chimera (c)  
 CC transducing DNA into cultured cells; and (d) harvesting the transduced  
 CC cells. The method is useful for ex vivo expansion of stem cells, and  
 CC enhancing the efficiency of the transduction of cultured stem cells by a  
 CC heterologous gene. The method is also useful for treating a patient  
 CC having a haematopoietic disorder. The expanded haematopoietic cells are  
 CC also useful in the treatment of cyclic neutropenia, aplastic anaemia,  
 CC thrombocytopenia, idiopathic neutropenia, Chediak-Higashi syndrome,  
 CC systemic lupus erythematosus (SLE), leukaemia, myelodysplastic syndrome  
 CC and myelofibrosis. The present sequence is an IL-3 mutant associated DNA  
 CC sequence. Note: The present sequence is included in the sequence listing  
 CC but is not mentioned anywhere else in the specification  
 XX

SQ Sequence 777 BP; 204 A; 227 C; 183 G; 163 T; 0 U; 0 Other;

## Alignment Scores:

Pred. No.: 1.38e-67 Length: 777  
 Score: 682.00 Matches: 130  
 Percent Similarity: 97.01% Conservative: 0  
 Best Local Similarity: 97.01% Mismatches: 4  
 Query Match: 89.15% Indels: 0  
 DB: 6 Gaps: 0

US-10-723-083-2 (1-142) x ABX00012 (1-777)

QY 9 SerGlyLeuGluGlyArgMetAlaProAlaArgSerProSerThrGlnProTyr 28  
 Db 376 TCTGGCGCGCGCTCCAACTGGACCGCTCGTTCCCGCTCCCGCTACCCAGCGGTGG 435  
 QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAla 48  
 Db 436 GAACAGTGAATGCCATCCAGAGAGGCCCGCGCTCTCTGAACCTGAGTAGAGACACTGCT 495  
 QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnGluProThr 68  
 Db 496 GCTGAGATGAATGAACAGTAGAAGTGAATCAAGAAATGTTTGCACCTCCAGAGCGCACT 555  
 QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88  
 Db 556 TGCCTACAGACCGCTGGAGCTGTACAGAGCGGCTCGGGGCGAGCCTCACCAAGCTC 615  
 QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProThrProGlu 108  
 Db 616 AAGGGCCCTTGACCATGATGGCCAGCCACTACAGCAGCACTGCCCTCCAAACCCCGAA 675  
 QY 109 ThrSerCysAlaThrGlnIleThrPheGluSerPheLysGluLeuLysAspPhe 128  
 Db 676 ACTTCTGTGCAACCCAGATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTC 735  
 QY 129 LeuLeuValIleProPheAspCysTyrGluProValGlnGlu 142  
 Db 736 CTGCTTGTATCCCTTTGACTGCTGGAGCCAGTCCAGGAG 777

## RESULT 4

ADJ14267

ID ADJ14267 standard; DNA; 777 BP.

XX AC ADJ14267;

XX DT 20-MAY-2004 (first entry)

XX DE DNA related to human interleukin-3 (IL-3) mutant protein - SEQ ID 55.

XX KW stem cell; antianaemic; immunostimulant; immunomodulator;

XX KW antiinflammatory; dermatological; immunosuppressive; cytostatic;

XX KW neuroprotective; haemopoietic disorder; gene therapy; myeloid; erythroid;

XX KW lymphoid; megakaryocyte; aplastic anaemia; periodic neutropenia;

XX KW Chediak-Higashi syndrome; systemic lupus erythematosus; leukaemia;

XX KW myelodysplastic syndrome; myelofibrosis; interleukin-3; IL-3; ds.

XX OS Unidentified.

XX PN US2003185790-A1.

XX PD 02-OCT-2003.

XX PF 26-FEB-2002; 2002US-00083446.

XX PR 24-NOV-1992; 92US-00981044.

XX PR 22-NOV-1993; 93WO-US011197.

XX PR 04-FEB-1994; 94US-00192325.

XX PR 02-FEB-1995; 95WO-US001185.

XX PR 06-APR-1995; 95US-00411795.

XX PR 06-JUN-1995; 95US-00446872.

XX PR 09-DEC-1996; 96US-00762227.

XX PA (BAUE/) BAUER S C.

XX PA (ABRA/) ABRAMS M A.

XX PA (BRAF/) BRAFORD-GOLDBERG S R.

XX PA (CAPA/) CAPARON M H.

XX PA (EAST/) EASTON A M.

XX PA (KLEI/) KLEIN B K.

XX PA (MCKE/) MCKEARN J P.

XX PA (OLIN/) OLINS P O.

XX PA (PAIK/) PAIK K.

XX PA (THOM/) THOMAS J W.

XX PI Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;

XX PI Klein BK, Mckearn JP, Olins PO, Paik K, Thomas JW;

XX WPI; 2004-096775/10.

XX Ex vivo expansion of stem cells, e.g. hematopoietic cells for treating

XX PT aplastic anemia, involves culturing the stem cells with growth medium

XX PT comprising chimera protein, and harvesting the cultured stem cells.

XX PS Disclosure; SEQ ID NO 55; 202pp; English.

XX CC The invention relates to a novel method whereby stem cells are ex vivo

XX CC expanded via culturing the stem cells with a growth medium comprising a

XX CC chimera protein, followed by harvesting of the cultured stem cells. The

XX CC method of the invention has antianaemic, immunostimulant,

XX CC immunomodulator, antiinflammatory, dermatological, immunosuppressive,

XX CC cytostatic and neuroprotective applications and may be useful to target

XX CC haemopoietic cells for gene therapy, preferably for treating patients

XX CC having a haemopoietic disorder characterised by decreased levels of

XX CC myeloid, erythroid, lymphoid, and/or megakaryocyte cells of haemopoietic

XX CC system. The expanded ex vivo cells may be used to treat neutropenia,

XX CC aplastic anaemia, periodic neutropenia, Chediak-Higashi syndrome,

XX CC systemic lupus erythematosus, leukaemia, myelodysplastic syndrome or

XX CC myelofibrosis. The current sequence is that of a DNA related to the human

XX CC interleukin-3 (IL-3) mutant protein of the invention.

XX SQ Sequence 777 BP; 204 A; 227 C; 183 G; 163 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 1.38e-67 Length: 777

Score: 682.00 Matches: 130

Percent Similarity: 97.01% Conservative: 0

Best Local Similarity: 97.01% Mismatches: 4

Query Match: 89.15% Indels: 0

DB: 12 Gaps: 0

US-10-723-083-2 (1-142) x ADJ14267 (1-777)

QY 9 SerGlyIleGluGlyArgMetAlaProAlaArgSerProSerProSerThrGlnProTrp 28

DB 376 TCTGGCGGGCTCCAAACATGGCAGCGCTCGTTCCCGCTCCCGTCTACCCAGCGGTGG 435

QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAla 48

DB 436 GAACAGTGAATGCCATCCAGGAGGCCCGCGCTCTCTGAACCTGAGTAGAGACTGCT 495

QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnProThr 68

DB 496 GCTGAGATGAATGAAACACGTAGAGTGATATCAGAAATGTTTGACCTCCAGAGCGGACT 555

QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88

DB 556 TGCCTACAGACCGCGCTGGAGCTGTACAAAGCAGGGCGCTCGGGGCGCAGCCTCACAAGCTC 615

QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProProThrProGlu 108

DB 616 AAGGGCCCTTGACCATGATGGCCAGCCACTAAAGCAGACTGCCCTCCAAACCCGGAA 675

QY 109 ThrSerCysAlaThrGlnIleIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128

DB 676 ACTTCTGTGCAACCCAGATTATCACCTTTGAAAGTTTCAAAGAGAACCTGAAGGACTTC 735

QY 129 LeuLeuValIleProPheAspCysTrpGluProValGlnGlu 142

DB 736 CTGCTGTGTCATCCCTTTGACTGTGGGAGCCAGTCCAGGAG 777

RESULT 5

AAT72724

ID AAT72724 standard; cDNA; 1610 BP.

XX AC AAT72724;

XX DT 17-SEP-1997 (first entry)

XX DE P53-GM-CSF immunostimulant fusion protein DNA.

XX KW P53-GM-CSF; granulocyte macrophage colony stimulating factor;

XX KW tumour suppressor gene; immunostimulant; cancer; therapy; ss.

XX OS Homo sapiens.

XX FH Key Location/Qualifiers

FT CDS 9..1610

FT /tag= a

FT /product= "p53-GM-CSF fusion protein"

FT mRNA 9..1187

FT /tag= b

FT /product= "p53"

FT mRNA 1188..1193

FT /tag= c

FT /product= "Ser-Arg linker"

FT mRNA 1194..1610

FT /tag= d

FT /product= "GM-CSF"

XX WO9724438-A1.

XX PN 10-JUL-1997.

XX PD 23-DEC-1996; 96WO-US020241.

XX PR 28-DEC-1995; 95US-00579823.

XX PA (ACTI-) ACTIVATED CELL THERAPY INC.

XX PI Laus R, Ruegg CL, Wu H;

XX



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DR WPI; 1997-363674/33.
XX P-PSDB; AAW19763.
PT Potent APC that activates T-cells to give multivalent cellular immune
PT response - can also induce a cytotoxic T-cell response in a vertebrate
XX subject.
XX
PS Example 7; Fig 11; 45pp; English.
XX
CC A nucleic acid molecule (AAAT72724) codes for a fusion protein (AAW19763)
CC comprising human p53 tumour suppressor protein and granulocyte-macrophage
CC colony stimulating factor (GM-CSF). It was prep'd. by PCR amplification of
CC p53 cDNA GM-CSF cDNA sequences (the GM-CSF antisense primer including a
CC hexahistidine tag sequence) and their fusion via a XbaI linker. Fusion
CC expression vectors can be used to transfect mammalian and insect cells.
CC The p53-GM-CSF fusion protein is used to generate anti-p53 immunity.
CC Tumour cells are eliminated by cytotoxic T lymphocytes activated in vivo
CC or in vitro by exposure to antigen-presenting cells exposed to the fusion
XX protein
XX
SQ Sequence 1610 BP; 383 A; 508 C; 407 G; 312 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.:      8,21e-67      Length:      1610
Score:          679.00      Matches:      128
Percent Similarity: 98.47%      Conservative: 1
Best Local Similarity: 97.71%      Mismatches: 2
Query Match:      88.76%      Indels:      0
DB:              2          Gaps:      0

US-10-723-083-2 (1-142) x AAT72724 (1-1610)
QY 12 GluGlyArgMetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisVal 31
DB 1185 GACTCTAGATCCGACCCCGCCGCTCGCCAGCCACAGCACAGCCCTGGGAGCATGTG 1244
QY 32 AsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMet 51
DB 1245 AATGCCATCCAGAGGCCGCGGCTCTCTGAACTGAGTAGACACACTGCTGTGATG 1304
QY 52 AsnGluThrValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGln 71
DB 1305 NATGAACAGTAGAGTATCTCAGAAATGTTTGACTCTCAGAGCGGACCTGCTTACAG 1364
QY 72 ThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyPro 91
DB 1365 ACCCGCTCGAGCTGTACAAGCAGGCGCTCGCGGCGAGCCTCACCAAGCTCAAGGGCCCC 1424
QY 92 LeuThrMetMetAlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCys 111
DB 1425 TTGACCATGATGCCAGCCCACTACAACAGCAGCACTGCCCTCCAAACCCCGGAAACTTCTCTGT 1484
QY 112 AlaThrGlnIleIleThrPheGluSerPheLeuGluAsnLeuLysAspPheLeuLeuVal 131
DB 1485 GCNAACCCAGATTATCACTTTGAAAGTTTCAAGAGAAACCTGAAGGACATTTCTGCTTGTG 1544
QY 132 IleProPheAspCysTrpGluProValGlnGlu 142
DB 1545 ATCCCTTTGACTGCTGGGAGCCAGCTCCAGAG 1577

RESULT 6
AAT34400
ID AAT34400 standard; DNA; 384 BP.
XX
AC AAT34400;
XX
XX
DT 25-MAR-2003 (revised)
DT 11-FEB-1997 (first entry)
XX
XX Granulocyte macrophage colony-stimulating factor coding sequence.
XX
XX GM-CSF; granulocyte macrophage colony-stimulating factor; expression;
KW construct; stable; production; ds.

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XX Homo sapiens.
XX
XX JP08173185-A.
PN
PD 09-JUL-1996.
XX
XX 28-APR-1987; 95JP-002633370.
XX
XX 28-APR-1987; 87JP-00106148.
XX
XX (AMGE-) AMGEN INC.
PA (KIRI ) KIRIN BREWERY KK.
PA
XX WPI; 1996-365600/37..
XX N-PSDB; AAW00103.
DR
XX Prodn. of human granulocyte macrophage colony-stimulating factor - by
XX culturing E. coli transformed with human GM-CSF DNA.
XX
XX Claim 1; Page 2; 16pp; Japanese.
XX
XX The present sequence encodes human granulocyte macrophage colony-
XX stimulating factor (hGM-CSF; n = 101, Ile). A series of oligonucleotides
XX were synthesised and ligated together to form a stable expression
XX construct. The technique is used for the efficient prodn. of a
XX glycoprotein with hGM-CSF activity. (Updated on 25-MAR-2003 to correct PF
XX field.)
XX
XX Sequence 384 BP; 101 A; 100 C; 86 G; 97 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.:      1.5e-67      Length:      384
Score:          678.00      Matches:      128
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match:      88.63%      Indels:      0
DB:              2          Gaps:      0

US-10-723-083-2 (1-142) x AAT34400 (1-384)
QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34
DB 1 ATGGCACCCAGCTCGATCACCGTCCGTCACCTCAACCATGGGAACATGTTAACGCAATC 60
QY 35 GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54
DB 61 CAGGAAGCTCGTCTGCTGTGTAACCTGTCTCGTACTGCTGCTGAAATGAACGAACT 120
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74
DB 121 GTTGAAGTGATCAGCGAAATGTTGATCTGCAGAACCGACTTGTCTGCAACCCGCTCTG 180
QY 75 GluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94
DB 181 GAACTGTACAAACAAGGCTGCGTGGTCTCTGACTAAACTGAAGGTCGCTGACTATG 240
QY 95 MetAlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGln 114
DB 241 ATGGCTAGCATTACAAACAGCATTTGTCGCCGACTCCGGAAACTTCTTGTGCTACTCAG 300
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134
DB 301 ATCATCACTTTCGAATCTTTCAAGAAACCTGAAAGATTTCTCTGCTGGTTATCCCGTTC 360
QY 135 AspCysTrpGluProValGlnGlu 142
DB 361 GATTGTGGGAACCGGTTTCAGGAA 384

RESULT 7
ABA96672
ID ABA96672 standard; cDNA; 384 BP.
XX

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AC ABA96672;  
 XX 23-APR-2002 (first entry)  
 XX Human granulocyte macrophage-colony stimulating factor (hGM-CSF) cDNA.  
 XX Human; GM-CSF; granulocyte macrophage-colony stimulating factor;  
 KW recombinant production; Escherichia coli; pET-11d vector; gene; ds.  
 XX Homo sapiens.  
 XX KR98053420-A.  
 XX 25-SEP-1998.  
 XX 26-DEC-1996; 96KR-00072523.  
 XX 26-DEC-1996; 96KR-00072523.  
 XX (CHEI-) CHEIL FOODS & CHEM INC.  
 XX Ahn DH, Lee HS, Mok H, Hah SH, Koh HG, Oh MS, Kim HS;  
 XX WPI; 1999-517067/43.  
 XX Human granulocyte macrophage-colony stimulating factor.  
 XX Example 1; Fig 4; 7pp; Korean.  
 XX The invention relates to the recombinant production of human granulocyte  
 CC macrophage-colony stimulating factor (hGM-CSF). NcoI and BamHI  
 CC restriction sites were inserted into hGM-CSF cDNA (ABA96672) via PCR  
 CC using primers ABA96670-ABA96671, prior to cleavage with these enzymes and  
 CC insertion into the pET-11d vector downstream of the T7/lac promoter to  
 CC form the plasmid pT7GMCSF. pT7GMCSF was transformed into Escherichia coli  
 CC BL21 (DE3) plys for recombinant expression of hGM-CSF. The present  
 CC sequence represents hGM-CSF cDNA  
 XX  
 SQ Sequence 384 BP; 93 A; 125 C; 97 G; 69 T; 0 U; 0 Other;  
 Alignment Scores:  
 Pred. No.: 15e-67 Length: 384  
 Score: 678.00 Matches: 128  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 88.63% Indels: 0  
 DB: 2 Gaps: 0  
 US-10-723-083-2 (1-142) x ABA96672 (1-384)  
 QY 15 MetAlaProAlaArgSerProSerThrGlnProThrGluHisValAlaAlaIle 34  
 DB 1 ATGGCACCCGCGCTCGCCAGCCCGCCAGCCGAGCCCTGGAGCATGTGATGCCATC 60  
 QY 35 GlnGluAlaArgGluLeuLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
 DB 61 CAGGAGGCGCGCTCTCTGAACCTGAGTAGACACTGCTGAGATGAATGAACA 120  
 QY 55 ValGluValIleSerGluMetPheAspLeuGlnProThrCysLeuGlnThrArgLeu 74  
 DB 121 GTAGAAGTCATCTCAGAAATGTTGACCTCCAGGAGCCGACCTGCCTCAGACCCGCTG 180  
 QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
 DB 181 GAGCTGTACAGCAGGCGCTCGGGGCGAGCCTCACCAGCTCAGGGCCCTTGACCATG 240  
 QY 95 MetAlaSerHisTyrlsGlnHisCysProThrProGluThrSerCysAlaThrGln 114  
 DB 241 ATGGCCAGGCACTACAGAGGAGCACTGCCCTCCAAACCCGGAACCTTCTGTGCAACCCAG 300  
 QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
 DB 301 AITATCACCTTTTGAAGATTTCAGAGAGAACCTGAGGACTTTTCTGCTTGTATCCCTTT 360

QY 135 AspCysTirpGluProValGlnGlu 142  
 DB 361 GACTGCTGGAGCCAGTCCAGGAG 384  
 RESULT 8  
 AAQ97208  
 ID AAQ97208 standard; DNA; 402 BP.  
 XX AC AAQ97208;  
 XX DT 25-AUG-1999 (first entry)  
 XX DE pMON13012 DNA sequence.  
 XX KW Interleukin; hIL-3; CSF; colony stimulating factor; cytokine; lymphokine;  
 XX mutant; mutein; fusion protein; linker; ss.  
 XX OS Synthetic.  
 XX OS Homo sapiens.  
 XX WO9521254-A1.  
 XX PD 10-AUG-1995.  
 XX PF 02-FEB-1995; 95WO-US001185.  
 XX PR 04-FEB-1994; 94US-00192325.  
 XX PA (SEAR) SEARLE & CO G D.  
 XX PI Bauer CS, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
 XX Klein BK, McKearn JP, Ollins PO, Paik K, Thomas JW;  
 XX WPI; 1995-283774/37.  
 XX P-PSDB; AAR79338.  
 XX Fusion proteins comprising a human interleukin-3 variant, a linker and  
 PT interleukin-3, a variant or a colony stimulating factor - useful to  
 PT increase haematopoietic cell prodn. in a mammal.  
 XX Example 19; Page 183; 447pp; English.  
 XX A new fusion protein is disclosed which has the formula R1-L-R2, R2-L-R1,  
 CC R1-R2, R2-R1, R1-L-R1 or R1-R1, where R1 is a mutant or variant of human  
 CC interleukin-3 (hIL-3), R2 is a second colony stimulating factor (CSF)  
 CC including cytokine, lymphokine, interleukin, haematopoietic growth factor  
 CC or IL-3 variant, and L is a linker. Generic sequences are described in  
 CC AAR03235 - AAW03242, and specifically claimed examples are shown in  
 CC AAR79298-R79335 and AAR79342-R79345. The fusion protein is made by  
 CC recombinant DNA techniques. Specifically claimed examples of DNA  
 CC sequences which encode these proteins are shown in AAQ97167-Q97204 and  
 CC AAQ97222-Q97227. The fusion protein is used to increase haematopoietic  
 CC cell production. It is also useful as an IL-3 antagonist or as a discrete  
 CC antigenic fragment for production of antibodies useful in immunoassays  
 CC and immunotherapy. Antagonists are used to block the growth of certain  
 CC cancer cells and in treatment of asthma. The fusion protein can also be  
 CC used to stimulate bone marrow and blood cell activation and growth in  
 CC vitro before infusion; and to treat diseases characterised by decreased  
 CC levels of myeloid, erythroid, lymphoid and/or megakaryocyte cells of the  
 CC haematopoietic system. The protein has the usual activity of both its  
 CC component proteins, but may have increased synergistic activity and  
 CC reduced undesired side effects  
 XX SQ Sequence 402 BP; 99 A; 124 C; 101 G; 78 T; 0 U; 0 Other;  
 Alignment Scores:  
 Pred. No.: 1.59e-67 Length: 402  
 Score: 678.00 Matches: 128  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 88.63% Indels: 0  
 DB: 2 Gaps: 0

US-10-723-083-2 (1-142) x AAQ97208 (1-402)

QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
 Db 1 ATGGCACCGGCTGTTCCCGTCCCGTCTACCCAGCGGTGGGAACACAGTGAATGCCATC 60

QY 35 GlnGluAlaArgGluLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
 Db 61 CAGGAGCGCGGGCTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAAGAACA 120

QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
 Db 121 GTAGAAGTGATATCAGAAATGTTTGAACCTCAGAGCGGCTGCTGCTGAGATGAAGAACA 180

QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
 Db 181 GAGCTGTACAGCAGGGCTGCGGGGAGCCCTCACCAGCTGCTGCTGAGATGAAGAACA 240

QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
 Db 241 ATGGCCAGCCACTACAAAGCAGCAGCTGCCCTCCAAACCCCGGAACTTCTGTGCAACCCAG 300

QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
 Db 301 ATTATCACCTTTGAAAGTGTTCAAAGAGAACCTGAGGACTTCTGCTGTGTCATCCCTTT 360

QY 135 AspCysTrpGluProValGlnGlu 142  
 Db 361 GACTGTGGGAGCCAGTCCAGGAG 384

RESULT 9  
 AAA03771  
 ID AAA03771 standard; DNA; 402 BP.  
 AC AAA03771;  
 DT 19-MAY-2000 (first entry)  
 DE Human G-CSF mutant DNA sequence SEQ ID NO:176.  
 KW Human; interleukin 3; IL-3; mutant; mutin; cytokine;  
 KW colony stimulating factor; haematopoietic growth factor; lymphokine;  
 KW fusion protein; haematopoietic disorder; infection; cancer;  
 KW radiation therapy; chemotherapy; bone marrow suppressive drug;  
 KW bone marrow activation; blood cell activation; blood transplant; ds.  
 OS Homo sapiens.  
 OS Synthetic.  
 PN US6022535-A.  
 PD 08-FEB-2000.  
 PF 06-JUN-1995; 95US-00469318.  
 PR 04-FEB-1994; 94US-00192325.  
 PR 02-FEB-1995; 95WO-US001185.  
 PR 06-APR-1995; 95US-00411795.  
 PA (SEAR ) SEARLE & CO G D.  
 PI Bauer SC, Abrams MA, Braford-Goldberg SR, Easton AM, Klein BK;  
 PI Paik K, Thomas JW, McKearn JP, Olins PO, Caparon MH;  
 XX WPI; 2000-160368/14.  
 XX Treating hematopoietic disorders with fusion proteins comprising mutated  
 XX interleukin-3 fused with secondary colony stimulating factors or other  
 XX interleukin-3 variants.  
 XX Example 19; Col 171-174; 276pp; English.

Methods have been developed for treating haematopoietic disorders with fusion proteins comprising recombinant, mutated human interleukin-3 (hIL-3) variants or mutant proteins (mutins) fused with secondary colony stimulating factors (CSFs) (e.g. cytokines, lymphokines, interleukin-3 and/or haematopoietic colony stimulating factors) or other interleukin-3 variants with or without a linker. The methods may be used in vivo to treat haematopoietic disorders resulting from bacterial, viral and fungal infections, cancer radiation therapy, chemotherapy or bone marrow suppressive drugs. They may also be used in vitro to stimulate bone marrow and blood cell activation and growth prior to infusion of the bone marrow and blood transplants into patients. IL-3 is a haematopoietic growth factor which has the property of being able to promote the survival, growth and differentiation of haematopoietic cells. The fusion molecules are characterised by possessing the usual activity of both of their constituent peptides and further by having a biological or physiological activity greater than the additive function of the IL-3 or second CSF alone (i.e. the peptides act synergistically). Their activity may also be further enhanced by the mutations they comprise. The variations may further reduce undesirable side effects associated with IL-3. AA03771 to AA03782 represent sequences used in the exemplification of the present invention

Sequence 402 BP; 99 A; 124 C; 101 G; 78 T; 0 U; 0 Other;

Alignment Scores:  
 Pred. No.: 1.59e-67 Length: 402  
 Score: 678.00 Matches: 128  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 88.63% Indels: 0  
 DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x AAA03771 (1-402)

QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
 Db 1 ATGGCACCGGCTGTTCCCGTCCCGTCTACCCAGCGGTGGGAACACAGTGAATGCCATC 60

QY 35 GlnGluAlaArgGluLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
 Db 61 CAGGAGCGCGGGCTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAAGAACA 120

QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
 Db 121 GTAGAAGTGATATCAGAAATGTTTGAACCTCAGAGCGGCTGCTGCTGAGATGAAGAACA 180

QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
 Db 181 GAGCTGTACAGCAGGGCTGCGGGGAGCCCTCACCAGCTGCTGCTGAGATGAAGAACA 240

QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
 Db 241 ATGGCCAGCCACTACAAAGCAGCAGCTGCCCTCCAAACCCCGGAACTTCTGTGCAACCCAG 300

QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
 Db 301 ATTATCACCTTTGAAAGTGTTCAAAGAGAACCTGAGGACTTCTGCTGTGTCATCCCTTT 360

QY 135 AspCysTrpGluProValGlnGlu 142  
 Db 361 GACTGTGGGAGCCAGTCCAGGAG 384

RESULT 10  
 ABX00086  
 ID ABX00086 standard; DNA; 402 BP.  
 XX ABX00086;  
 AC ABX00086;  
 XX 18-DEC-2002 (first entry)  
 DT Human interleukin-3 associated DNA sequence #77.  
 XX Haematopoietic factor; GM-CSF; colony stimulating factor; CSF-1; ds;  
 KW

G-CSF; G-CSFser17; c-mpl ligand; TPO; MGDF; erythropoietin; flt3 ligand; human growth hormone; B-cell growth factor; leukaemia; B-cell differentiation factor; eosinophil differentiation factor; stem cell factor; SCF; cyclic neutropenia; aplastic anaemia; thrombocytopenia; idiopathic neutropenia; Chediak-Higashi syndrome; systemic lupus erythematosus; SLE; myelodysplastic syndrome; myelofibrosis; Interleukin-3; IL-3; stem cell.

Unidentified.

US6436387-B1.

20-AUG-2002.

09-DEC-1996; 96US-00762227.

24-NOV-1992; 92US-00981044.

22-NOV-1993; 93WO-US011197.

04-FEB-1994; 94US-00192325.

06-FEB-1995; 95WO-US001185.

06-APR-1995; 95US-00411795.

06-JUN-1995; 95US-00446872.

(SEAR ) SEARLE & CO G D.

Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;

Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;

WPI; 2002-749206/81.

Ex vivo expansion of stem cells, for enhancing transduction efficiency of

cultured stem cells, comprises culturing stem cells in growth medium

having mutant interleukin-3, and hematopoietic factor, and harvesting

cultured cells.

Disclosure; Col 299-300; 203pp; English.

The invention relates to ex vivo expansion of stem cells, comprises

culturing stem cells with a growth medium comprising a chimera protein,

and harvesting the cultured stem cells. The chimera is based on a

mutated human interleukin-3 (IL-3) sequence coupled to a hematopoietic

factor (e.g. GM-CSF (colony stimulating factor), CSF-1, G-CSF, G-

CSFser17, c-mpl ligand TPO, MGDF, erythropoietin, IL-1-13, IL-15, IL-16,

flt3 ligand, human growth hormone, B-cell growth factor, B-cell

differentiation factor, eosinophil differentiation factor and stem cell

factor (SCF)) via a peptide linker. The formula for the chimera is given

in the specification. Also included is a method for enhancing the

efficiency of the transduction of cultured stem cells by a heterologous

gene, comprising: (a) removing stem cells from a patient or donor; (b)

culturing the stem cells with a growth medium comprising the chimera (c)

transducing DNA into cultured cells; and (d) harvesting the transduced

cells. The method is useful for ex vivo expansion of stem cells, and

enhancing the efficiency of the transduction of cultured stem cells by a

heterologous gene. The method is also useful for treating a patient

having a hematopoietic disorder. The expanded hematopoietic cells are

also useful in the treatment of cyclic neutropenia, aplastic anaemia,

thrombocytopenia, idiopathic neutropenia, Chediak-Higashi syndrome,

systemic lupus erythematosus (SLE), leukaemia, myelodysplastic syndrome

and myelofibrosis. The present sequence is an IL-3 mutant associated DNA

sequence. Note: The present sequence is included in the sequence listing

but is not mentioned anywhere else in the specification

XX Sequence 402 BP; 99 A; 124 C; 101 G; 78 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 1,596-67 Length: 402

Score: 678.00 Matches: 129

Percent Similarity: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0

Query Match: 88.63% Indels: 0

DB: 6 Gaps: 0

US-10-723-083-2 (1-142) x ABX00086 (1-402)

QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTirpGluHisValaenAlaile 34  
 Db 1 ATGGCACCGGCTCTTCCCGTCCCGTCTACCCAGCGCTGGGAACACGCTGAATGCCATC 60  
 QY 35 GlnGluAlaArgArgLeuLeuLeuSerArgAspThrAlaAlaGluMetAenGluThr 54  
 Db 61 CAGGAGCGCGCGTCTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAATGAACA 120  
 QY 55 ValGluValileSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
 Db 121 GTAGAAGTGATATCAGAAATGTTGACCTCCAGGAGCCGACTTGCCTCAGACCCCGCTG 180  
 QY 75 GluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
 Db 181 GAGCTGTACAAGCAGGGCTTGGGGGCGAGCTTCAACAGCTCAAGGGCCCTTGACCATG 240  
 QY 95 MetAlaSerHisTyrLysGlnHisCysProThrProGluThrSerCysAlaThrGln 114  
 Db 241 ATGGCCAGGCACCTACAAAGCAGCACTGCCCTCCAAACCCGGAACCTTCTGTGCAACCCAG 300  
 QY 115 IleIleThrPheGluSerPheLysGluAenLeuLysAspPheLeuValleProPhe 134  
 Db 301 ATTATCACCTTTGAAAGTTTCAAGAGAACCTTGAAGGAGCTTCTGCTTGTGTCATCCCTTT 360  
 QY 135 AspCysTirpGluProValGlnGlu 142  
 Db 361 GACTGCTGGAGCCAGTCCAGGAG 384

RESULT 11

ADJ14388

ID ADJ14388 standard; DNA; 402 BP.

AC ADJ14388;

XX 20-MAY-2004 (first entry)

XX DNA related to human interleukin-3 (IL-3) mutant protein - SEQ ID 151.

XX stem cell; antianaemic; immunostimulant; immunomodulator;  
 XX antiinflammatory; dermatologic; immunosuppressive; cytostatic;  
 XX neuroprotective; haemopoietic disorder; gene therapy; myeloid; erythroid;  
 XX lymphoid; megakaryocyte; aplastic anaemia; periodic neutropenia;  
 XX Chediak-Higashi syndrome; systemic lupus erythematosus; leukaemia;  
 XX myelodysplastic syndrome; myelofibrosis; interleukin-3; IL-3; ds.

XX Unidentified.

XX US2003185790-A1.

XX 02-OCT-2003.

XX 26-FEB-2002; 2002US-00083446.

XX 24-NOV-1992; 92US-00981044.

XX 22-NOV-1993; 93WO-US011197.

XX 04-FEB-1994; 94US-00192325.

XX 02-FEB-1995; 95WO-US001185.

XX 06-APR-1995; 95US-00411795.

XX 06-JUN-1995; 95US-00446872.

XX 09-DEC-1996; 96US-00762227.

XX (BAUE/) BAUER S C.

XX (ABRA/) ABRAMS M A.

XX (BRAD/) BRAFORD-GOLDBERG S R.

XX (CAPA/) CAPARON M H.

XX (EAST/) EASTON A M.

XX (KLEI/) KLEIN B K.

XX (MCKE/) MCKEARN J P.

XX (OLIN/) OLINS P O.

XX (PAIK/) PAIK K.

XX (THOM/) THOMAS J W.

XX

PI Bauer SC, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
 PI Klein BK, Mckearn JP, Olines PO, Paik K, Thomas JW;  
 DR WPI; 2004-096775/10.  
 XX  
 XX Ex vivo expansion of stem cells, e.g. hematopoietic cells for treating  
 PT aplastic anemia, involves culturing the stem cells with growth medium  
 PT comprising chimera protein, and harvesting the cultured stem cells.  
 XX  
 PS Disclosure; SEQ ID NO 176; 202pp; English.  
 XX  
 XX The invention relates to a novel method whereby stem cells are ex vivo  
 CC expanded via culturing the stem cells with a growth medium comprising a  
 CC chimera protein, followed by harvesting of the cultured stem cells. The  
 CC method of the invention has antianemic, immunostimulant,  
 CC immunomodulatory, antiinflammatory, dermatological, immunosuppressive,  
 CC cytostatic and neuroprotective applications and may be useful to target  
 CC hematopoietic cells for gene therapy, preferably for treating patients  
 CC having a haemopoietic disorder characterised by decreased levels of  
 CC myeloid, erythroid, lymphoid, and/or megakaryocyte cells of haemopoietic  
 CC system. The expanded ex vivo cells may be used to treat neutropenia,  
 CC aplastic anaemia, periodic neutropenia, Chediak-Higashi syndrome,  
 CC systemic lupus erythematosus, leukaemia, myelodysplastic syndrome or  
 CC myelofibrosis. The current sequence is that of a DNA related to the human  
 CC interleukin-3 (IL-3) mutant protein of the invention.  
 XX  
 SQ Sequence 402 BP; 99 A; 124 C; 101 G; 78 T; 0 U; 0 Other;

Alignment Scores:  
 Pred. No.: 1.59e-67 Length: 402  
 Score: 678.00 Matches: 128  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 88.63% Indels: 0  
 DB: 12 Gaps: 0

US-10-723-083-2 (1-142) x ADJ14388 (1-402)

QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
 DB 1 ATGGCACCGGCTCGTTCCCGTCCCGTCTACCCAGCGGTGGGAACACAGTGAATGCCATC 60  
 QY 35 GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
 DB 61 CAGGAGGCGCGGCTCTCTGAAACCTGAGTAGACACTGCTGCTGAGATGAATGAACA 120  
 QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
 DB 121 GTAGAAGTGAATATCAGAAATGTTTGACCTCCAGAGCCGACTTGCCTACAGACCCGCTG 180  
 QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
 DB 181 GAGCTGTACAAAGCAGGCGCTCGGGGCGAGCTCAACCAAGCTCAAGGGCCCTTGACCATG 240  
 QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
 DB 241 ATGGCCAGGCACATACAAAGCAGCAGCTGCCCTCCAAACCCCGGAAACTTCTGTGTGCAACCCAG 300  
 QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
 DB 301 ATTATCACCTTTGAAGATTTCAAAGAGAACCTGAGGACTTTCCTGTTGTATCCCTTT 360  
 QY 135 AspCysTrpGluProValGlnGlu 142  
 DB 361 GACTGTGGGAGCCAGTCCAGGAG 384  
 RESULT 12  
 AAN81322  
 ID AAN81322 standard; DNA; 410 BP.  
 XX  
 AC AAN81322;  
 XX  
 DT 25-MAR-2003 (revised)

DT 06-JAN-1991 (first entry)  
 XX  
 DE Sequence (I) encoding human granulocyte macrophage colony stimulating  
 DE factor (hGMCSF) with Ile at 101.  
 XX  
 DE Leukopenia therapy; bone marrow transplant; ds.  
 KW  
 XX Homo sapiens.  
 OS  
 XX  
 FH Key Location/Qualifiers  
 FT misc\_feature complement(1..4)  
 FT /\*tag= a  
 FT /note= "Sticky end"  
 FT misc\_feature complement(407..410)  
 FT /\*tag= b  
 FT /note= "Sticky end"  
 FT  
 XX JP63269983-A.  
 PN 08-NOV-1988.  
 XX  
 PD 28-APR-1987; 87JP-00106148.  
 XX  
 PF 28-APR-1987; 87JP-00106148.  
 XX  
 PR 28-APR-1987; 87JP-00106148.  
 XX  
 XX (AMGE-) AMGEN.  
 PA (KIRI ) KIRIN BREWERY KK.  
 PA  
 XX WPI; 1988-358227/50.  
 DR  
 XX Human granulocyte macrophage colony stimulus factor prepn. - by forming  
 PT plasmid contg. genetic information and Escherichia coli with the plasmid.  
 XX  
 PS Disclosure; Fig 7(1), Page 488; 17pp; Japanese.  
 XX  
 CC An expression plasmid contg. the DNA is prep'd, and used to transform  
 CC E.coli. The transformant is cultured and hGMCSF is produced into the  
 CC medium. hGMCSF stimulates the granulocyte macrophage system stem cell and  
 CC promotes the formation of the granulocyte macrophage. hGMCSF is used in  
 CC curing leukopenia caused by radiation therapy, or chemical therapy, or  
 CC in promptly propagating leukocytes after bone marrow transplantation.  
 CC (Updated on 25-MAR-2003 to correct PA field.)  
 XX  
 SQ Sequence 410 BP; 109 A; 106 C; 91 G; 104 T; 0 U; 0 Other;

Alignment Scores:  
 Pred. No.: 1.64e-67 Length: 410  
 Score: 678.00 Matches: 128  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 88.63% Indels: 0  
 DB: 1 Gaps: 0

US-10-723-083-2 (1-142) x AAN81322 (1-410)

QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
 DB 11 ATGGCACCGCTCGATCAGCTCCCGTCCACTCAACCATGGGAACATGTTAACGCAATC 70  
 QY 35 GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
 DB 71 CAGGAAGCTCGCTGCTGCTGAAACCTGCTCGTGATGCTGCTGCTGAAATGAACGAAACT 130  
 QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
 DB 131 GTTGAAGTGAATCAGCAAAATGTTGATCTGCAGAACCCGACTTGTCTGCAAAACCCGCTG 190  
 QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
 DB 191 GAACTGTACAAACAGGTCTGCGTGGTTCTCTGACTAAACTGAAGAGTCCGCTGACTATG 250  
 QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114

Db 251 ATGGTAGCCATTACAAACAGCAATGTCCGCCGACTCCGAAACTTCTTGTGCTACTCAG 310  
 QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPhe 134  
 Db 311 ATCATACATTTCGAATCTTTCAAGAAACCTGAAAGATTTCCTGCTGGTTATCCCGTTC 370  
 QY 135 AspCysTrpGluProValGlnGlu 142  
 Db 371 GATTGTTGGGAACCGGTTTCAGGAA 394

RESULT 13  
 AAN90383  
 ID AAN90383 standard; DNA; 415 BP.  
 AC AAN90383;  
 XX  
 DT 25-MAR-2003 (revised)  
 DT 01-NOV-1989 (first entry)  
 XX  
 DE Synthetic gene for human granulocyte colony stimulating factor.  
 XX  
 KW DNA; BspMI; restriction sites; blunt ends; fusion proteins;  
 KW synthetic gene; human granulocyte colony stimulating factor.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 XX  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 14..397  
 FT /\*tag= a  
 XX  
 XX GB2212160-A.  
 PN 19-JUL-1989.  
 XX  
 XX  
 PF 13-NOV-1987; 87GB-00726581.  
 XX  
 PR 13-NOV-1987; 87GB-00026581.  
 XX  
 PA (BRBI-) BRITISH BIO-TECHN L.  
 XX  
 PI Edwards RM;  
 XX  
 XX WPI; 1989-208959/29.  
 DR P-PSDB; AAP90118.  
 XX  
 XX DNA including recognition site for BspMI enzyme - allowing generation of  
 PT blunt end for fusion in prodn. of fusion proteins.  
 XX  
 PS Disclosure; Fig 2; 23pp; English.  
 XX  
 CC Synthetic gene for human granulocyte colony stimulating factor (GM-CSF)  
 CC contg. useful restriction sites, and a BspMI site. See corresp. AAP90118.  
 CC Useful for generating blunt ends in fusion protein prodn. (Updated on 25-  
 CC MAR-2003 to correct PF field.) (Updated on 25-MAR-2003 to correct PR  
 CC field.) (Updated on 25-MAR-2003 to correct PA field.)  
 XX  
 XX Sequence 415 BP; 104 A; 130 C; 104 G; 77 T; 0 U; 0 Other;  
 SQ

Alignment Scores:  
 Pred. No.: 1.67e-67 Length: 415  
 Score: 678.00 Matches: 128  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 88.63% Indels: 0  
 DB: 1 Gaps: 0

US-10-723-083-2 (1-142) x AAN90383 (1-415)

QY 15 MetaAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
 Db 14 ATGGCACCGCCCGGTTCACCCAGCCCCAGCAGCAGCCCTGGGAGCATGTGATGCCATC 73

QY 35 GlnGluAlaArgGluLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
 Db 74 CAGGAGCCCGCGGTCTCTCTGAACCTGAGTACAGACACTGCTGCTGAGATGAATGAACA 133  
 QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
 Db 134 GTAGAAGTGATATCAGAAATGTTTGACCTCCAGGAGCCGACTTGCCTACAGACCCGCTG 193  
 QY 75 GluLeuTyrlsGlnGluLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
 Db 194 GAGCTGTACAAGCAGGGGCTGCGGGGCGACCTCACCAAGCTCAAGGGCCCCCTGACCATG 253  
 QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
 Db 254 ATGGCCAGCCACTACAAGCAGCAGCTGCCCTCCACCCCGGAACCTCTGTGCAACCCAG 313  
 QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPhe 134  
 Db 314 ATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAAGGACTTCTGCTTGTCTATCCCTTT 373  
 QY 135 AspCysTrpGluProValGlnGlu 142  
 Db 374 GACTGCTGGAGCCAGTCCAGGAG 397

RESULT 14  
 AAN90274  
 ID AAN90274 standard; DNA; 415 BP.  
 AC AAN90274;  
 XX  
 DT 25-MAR-2003 (revised)  
 DT 01-NOV-1989 (first entry)  
 XX  
 DE Synthetic human granulocyte colony stimulating factor.  
 XX  
 KW Synthetic DNA; human granulocyte colony stimulating factor;  
 KW restriction sites; cassette mutagenesis; expression system.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 14..397  
 FT /\*tag= a  
 XX  
 XX GB2212159-A.  
 PN 19-JUL-1989.  
 XX  
 PF 13-NOV-1987; 87GB-00026580.  
 XX  
 PR 13-NOV-1987; 87GB-00026580.  
 XX  
 PA (BRBI-) BRITISH BIO-TECHN L.  
 XX  
 PI Edwards RM;  
 XX  
 XX WPI; 1989-208958/29.  
 DR P-PSDB; AAP90115.  
 XX  
 XX Human granulocyte-macrophage colony stimulating factor - synthetic DNA  
 PT includes restriction sites for cassette mutagenesis and incorporation in  
 PT expression systems.  
 XX  
 PS Claim 2; Page 14 and fig 3a; 21pp; English.  
 XX  
 CC Synthetic DNA encoding human granulocyte colony stimulating factor (GM-  
 CC CSF), see corresp. AAP90115. Has useful restriction sites for: HindIII;  
 CC BspMI; NcoI; BstRII; BsmI; EcoRV; BglI; ApaI; BclI; XbaI; BamHI; and  
 CC EcoRI. Used to facilitate cassette mutagenesis of selected regions.  
 CC Synthesised by phosphoramidite chemistry, by dividing desired gene into  
 CC 18 oligomers. (Updated on 25-MAR-2003 to correct PA field.)  
 XX

SQ Sequence 415 BP; 104 A; 130 C; 104 G; 77 T; 0 U; 0 Other;

# Alignment Scores:

Pred. No.: 1.67e-67 Length: 415  
 Score: 678.00 Matches: 128  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 88.63% Indels: 0  
 DB: 1 Gaps: 0

US-10-723-083-2 (1-142) x AAN90274 (1-415)

QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
 DB 14 ATGGCACCAGCCCGGCTCTCCAGACCTGAGTAGAGACACTGCTGCTGAGATGAATGAACA 73  
 QY 35 GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
 DB 74 CAGGAGGCCCGGGCTCTCCAGACCTGAGTAGAGACACTGCTGCTGAGATGAATGAACA 133  
 QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
 DB 134 GTAGAAGTGATATCAGAAATGTTTGACCTCCAGAGCGGACTTGCCTACAGACCCGCTG 193  
 QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrIlysLeuLysGlyProLeuThrMet 94  
 DB 194 GAGCTGTACAAGCAGGGCGCTGCGGGGAGCCCTCACCAGCTCAAGGGCCCTTGACCATG 253  
 QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
 DB 254 ATGGCCAGCACCTACAGCAGCAGCTGCCCTCCAAACCCCGGAAACTTCTGTGCAACCCAG 313  
 QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
 DB 314 ATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTCTGCTTGTGTCATCCCTTT 373  
 QY 135 AspCysTrpGluProValGlnGlu 142  
 DB 374 GACTGTGGAGCCAGTCCAGGAG 397

# RESULT 15

ID AAQ97183  
 AC AAQ97183 standard; DNA; 822 BP.  
 AC AAQ97183;  
 XX  
 XX  
 XX  
 DT 25-AUG-1999 (first entry)  
 XX  
 DE pMON13035 DNA encoding IL-3 fusion protein.  
 XX  
 XW Interleukin; hIL-3; CSF; colony stimulating factor; cytokine; lymphokine;  
 XW mutant; mutein; fusion protein; linker; ss.  
 XX  
 OS Synthetic.  
 OS Homo sapiens.  
 XX  
 XX W09521254-A1.  
 XX  
 XX 10-AUG-1995.  
 XX  
 PF 02-FEB-1995; 95WO-US0001185.  
 XX  
 PR 04-FEB-1994; 94US-00192325.  
 XX  
 XX (SEARLE & CO G D.  
 PA  
 XX  
 XX Bauer CS, Abrams MA, Braford-Goldberg SR, Caparon MH, Easton AM;  
 PI Klein BK, McKearn JP, Olins PO, Paik K, Thomas JW;  
 XX  
 XX WPI; 1995-283774/37.  
 DR P-PSDB; AAR79320.  
 XX  
 PT Fusion proteins comprising a human interleukin-3 variant, a linker and

PT interleukin-3, a variant or a colony stimulating factor - useful to  
 PT increase haematopoietic cell prodn. in a mammal.

Claim 22; Page 167-168; 447pp; English.

CC A new fusion protein is disclosed which has the formula R1-L-R2, R2-L-R1,  
 CC R1-R2, R2-R1, R1-L-R1 or R1-R1, where R1 is a mutant or variant of human  
 CC interleukin-3 (hIL-3), R2 is a second colony stimulating factor (CSF)  
 CC including cytokine, lymphokine, interleukin, haematopoietic growth factor  
 CC or IL-3 variant, and L is a linker. Genetic sequences are described in  
 CC AAR03235 - AAW03242, and specifically claimed examples are shown in  
 CC AAR79298-R79335 and AAR79342-R79345. The fusion protein is made by  
 CC recombinant DNA techniques. Specifically claimed examples of DNA  
 CC sequences (including the present sequence) which encode these proteins  
 CC are shown in AAQ97167-Q97204 and AAQ97222-Q97227. The fusion protein is  
 CC used to increase haematopoietic cell production. It is also useful as an  
 CC IL-3 antagonist or as a discrete antigenic fragment for production of  
 CC antibodies useful in immunoassays and immunotherapy. Antagonists are used  
 CC to block the growth of certain cancer cells and in treatment of asthma.  
 CC The fusion protein can also be used to stimulate bone marrow and blood  
 CC cell activation and growth in vitro before infusion; and to treat  
 CC diseases characterised by decreased levels of myeloid, erythroid,  
 CC lymphoid and/or megakaryocyte cells of the haematopoietic system. The  
 CC protein has the usual activity of both its component proteins, but may  
 CC have increased synergistic activity and reduced undesired side effects  
 XX  
 SQ Sequence 822 BP; 222 A; 244 C; 180 G; 176 T; 0 U; 0 Other;

# Alignment Scores:

Pred. No.: 4.25e-67 Length: 822  
 Score: 678.00 Matches: 128  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 88.63% Indels: 0  
 DB: 2 Gaps: 0

US-10-723-083-2 (1-142) x AAQ97183 (1-822)

QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
 DB 439 ATGGCACCAGCCCGGCTCTCCAGACCTGAGTAGAGACACTGCTGCTGAGATGAACA 498  
 QY 35 GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
 DB 499 CAGGAGGCCCGGGCTCTCCAGACCTGAGTAGAGACACTGCTGCTGAGATGAACA 558  
 QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
 DB 559 GTAGAAGTGATATCAGAAATGTTTGACCTCCAGAGCCGACTTGCCTACAGACCCGCTG 618  
 QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
 DB 619 GAGCTGTACAAGCAGGGCGCTGCGGGGAGCCCTCACCAGCTCAAGGGCCCTTGACCATG 678  
 QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
 DB 679 ATGGCCAGCACCTACAGAGCAGCACTGCCCTCCAAACCCCGGAAACTTCTGTGCAACCCAG 738  
 QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
 DB 739 ATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTCTGCTTGTGTCATCCCTTT 798  
 QY 135 AspCysTrpGluProValGlnGlu 142  
 DB 799 GACTGTGGAGCCAGTCCAGGAG 822

Search completed: March 11, 2005, 18:36:51  
 Job time : 480 secs





GenCore version 5.1.6  
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OM protein - nucleic search, using frame\_plus\_p2n model

Run on: March 11, 2005, 17:26:51 ; Search time 3628 Seconds  
(without alignments)  
1896.538 Million cell updates/sec

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Perfect score: 765  
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Ygapop 10.0 , Ygapext 0.5  
Fgapop 6.0 , Fgapext 7.0  
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Searched: 4708233 seqs, 24227607955 residues  
Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0  
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Post-processing: Minimum Match 0%  
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Listing first 45 summaries

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-UNITS=bits -START=1 -END=1 -MATRIX=blom62 -TRANS=human40.cdi -LIST=45  
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2: gb.htg.\*  
3: gb.in.\*  
4: gb.om.\*  
5: gb.ov.\*  
6: gb.pat.\*  
7: gb.ph.\*  
8: gb.pl.\*  
9: gb.pr.\*  
10: gb.ro.\*  
11: gb.sts.\*  
12: gb.sy.\*  
13: gb.un.\*  
14: gb.vi.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	682	89.2	777	6 AR202206	AR202206 Sequence
2	682	89.2	777	6 AR223208	AR223208 Sequence
3	678	88.6	384	6 E11629	E11629 hGM-CSF-enc
4	678	88.6	402	6 AR202280	AR202280 Sequence

5	678	88.6	402	6 AR223282	AR223282 Sequence
6	678	88.6	415	6 A00367	A00367 Artificial
c 7	678	88.6	415	6 A00368	A00368 Artificial
8	678	88.6	415	6 A14305	A14305 GM-CSF gene
c 9	678	88.6	415	6 A14306	A14306 GM-CSF gene
10	678	88.6	822	6 AR202220	AR202220 Sequence
11	678	88.6	822	6 AR223222	AR223222 Sequence
12	678	88.6	903	6 AR202217	AR202217 Sequence
13	678	88.6	903	6 AR223219	AR223219 Sequence
14	676	88.4	1630	6 BD222938	BD222938 Heteromim
15	676	88.4	1630	6 AX023365	AX023365 Sequence
16	673	88.0	384	6 A20082	A20082 Pcti-Hind I
17	673	88.0	384	6 E11628	E11628 hGM-CSF-enc
c 18	673	88.0	392	6 A20083	A20083 Pcti-Hind I
19	673	88.0	392	6 I49837	I49837 Sequence 1
20	673	88.0	418	6 A11762	A11762 Artificial
c 21	673	88.0	418	6 A11763	A11763 Artificial
22	673	88.0	435	6 CQ834914	CQ834914 Sequence
23	673	88.0	435	6 CQ834915	CQ834915 Sequence
24	673	88.0	435	6 AR533388	AR533388 Sequence
25	673	88.0	435	6 BD105958	BD105958 Animal mo
26	673	88.0	436	6 E01740	E01740 gene encodi
27	673	88.0	505	6 I08646	I08646 Sequence 3
28	673	88.0	505	9 HUMCSFGMB	M11734 Human granu
29	673	88.0	644	6 E01141	E01141 cDNA encodi
30	673	88.0	660	6 AR364645	AR364645 Sequence
31	673	88.0	661	6 E00951	E00951 cDNA encodi
32	673	88.0	661	6 I04859	I04859 Sequence 3
33	673	88.0	661	6 AR363793	AR363793 Sequence
34	673	88.0	756	6 CQ721607	CQ721607 Sequence
35	673	88.0	756	6 CQ803372	CQ803372 Sequence
36	673	88.0	763	6 E02287	E02287 DNA encodin
37	673	88.0	787	6 I08093	I08093 Sequence 1
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39	673	88.0	787	6 I08401	I08401 Sequence 1
40	673	88.0	787	6 I09140	I09140 Sequence 1
41	673	88.0	787	6 I09160	I09160 Sequence 21
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ALIGNMENTS

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LOCUS AR202206 777 bp DNA linear PAT 20-APR-2002  
DEFINITION Sequence 55 from patent US 6361977.  
ACCESSION AR202206  
VERSION AR202206.1 GI:20256745  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE 1 (bases 1 to 777)  
AUTHORS Bauer, S. Christopher., Abrams, M. Allen., Braford-Goldberg, S. Ruth., Caparon, M. Helena., Easton, A. Michael., Klein, B. Kure., McKearn, J. P., Olin, P. O., Paik, K. and Thomas, J. W.  
TITLE Methods of using multivalent IL-3 hematopoiesis fusion protein  
JOURNAL Patent: US 6361977-A 55 26-MAR-2002;  
FEATURES Location/Qualifiers  
source 1..777  
/organism="unknown"  
/mol\_type="unassigned DNA"

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Best Local Similarity: 97.01% Mismatches: 4  
Query Match: 89.15% Indels: 0

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DB: 6 Gaps: 0
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QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAla 48
Db 436 GAACAGTGAATGCCATCCAGAGGCGCGGCTCTCTGACCTGATAGACACTGCT 495
QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnProTrp 68
Db 496 GCTCAGATGAATGAACAGTAGAAGTATCAGAAATGTTTGACCTCCAGGCGGACT 555
QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88
Db 556 TGCTACAGACCGCTCGAGCTGTACAGCAGGCGCTCGCGGCGGAGCTCCCAAGCTC 615
QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProProThrProGlu 108
Db 616 AAGGCGCCCTTGACCATGATGCCAGCCACTACAAGCAGACTGCCCTCCAAACCCGGAA 675
QY 109 ThrSerCysAlaThrGlnIleIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128
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QY 129 LeuLeuValIleProPheAspCysTrpGluProValGlnGlu 142
Db 736 CTGCTGTATCCCTTTGACTCTGGAGCCAGTCCAGAG 777

RESULT 3
E11629
LOCUS E11629 384 bp DNA linear PAT 29-SEP-1997
DEFINITION hgm-CSF-encoding DNA for efficient expression in E.coli.
ACCESSION E11629
VERSION E11629.1 GI:22025265
KEYWORDS JP 1996173185-A/2.
SOURCE unidentified
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 384)
AUTHORS Matsuki,S., Ozawa,T. and Tamai,Y.
TITLE PRODUCTION OF STIMULATING FACTOR FOR HUMAN GRANULOCYTIC MACROPHAGE
COLONY
JOURNAL Patent: JP 1996173185-A 2 09-JUL-1996;
COMMENT AMGEN, KIRIN BREWERY CO LTD
OS None
OC Artificial sequences.
PN JP 1996173185-A/2
PD 09-JUL-1996
PF 28-APR-1987 JP 1995263370
PI MATSUKI SHIGERU, OZAWA TADASHI, TAMAI YUKIO
PC C12P21/02,C12N1/21,C12N15/09,(C12P21/02,C12R1:19),(C12N1/21,
CC strandedness: Double;
CC topology: Linear;
CC Key Location/Qualifiers
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Query Match: 88.63% Indels: 0
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Db 61 CAGGAAGCTCGTCTGCTGAACCTGCTCTCGTACTGCTGCTGAATGAACGAACT 120  
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Db 121 GTTGAAGTGTATCAGCGAAATGTTTCGATCTGCAGGAACCGACTTGTCTGCAACCCGCTCG 180  
Qy 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 181 GAACCTGACAAACAAGGTCTGCGTGTCTCTGACTAAACTGAAAGGTCCGCTGACTATG 240  
Qy 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
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RESULT 4  
LOCUS AR202280 402 bp DNA linear PAT 20-APR-2002  
DEFINITION Sequence 176 from patent US 6361977.  
ACCESSION AR202280  
VERSION AR202280.1 GI:20256819  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 402)  
AUTHORS Bauer, S., Christopher., Abrams, M., Allen., Braford-Goldberg, S., Ruth., Caparon, M., Helena., Easton, A., Michael., Klein, B., Kure., McKearn, J. P., Oling, P. O., Paik, K. and Thomas, J. W.  
TITLE Methods of using multivariant IL-3 hematopoiesis fusion protein  
JOURNAL Patent: US 6361977-A 176 26-MAR-2002;  
FEATURES Location/Qualifiers  
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Score: 678.00 Matches: 128  
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DB: 6 Gaps: 0  
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Qy 15 MetAlaProAlaArgSerProSerProSerThrGlnProThrGluHisValAsnAlaIle 34  
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Qy 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
Db 121 GTAGAAGTGTATCAGAAATGTTGACCTCCAGAGCCGACTTGCCTACAGACCCGCTG 180  
Qy 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 181 GAGCTGTACAAAGCAGGCGCTTCCCGGCGAGCCTCACCAGCTCAAGGCGCCCTTGACCATG 240  
Qy 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
Db 241 ATGGCCAGCCACTTACAAAGCAGCACTGCCCTCCAAACCCCGGAAACTTCTCTGTGCAACCCAG 300  
Qy 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPhe 134  
Db 301 ATATCACCTTTGAAAGTTTCAAGAGAACCTGGAAGACTTCTGCTGTGTCATCCCTTT 360  
Qy 135 AspCysTrpGluProValGlnGlu 142  
Db 361 GACTGCTGGAGCCAGTCCAGGAG 384  
RESULT 6  
LOCUS A00367 415 bp DNA linear PAT 09-MAR-1993

Qy 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPhe 134  
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Qy 135 AspCysTrpGluProValGlnGlu 142  
Db 361 GACTGCTGGAGCCAGTCCAGGAG 384  
RESULT 5  
LOCUS AR223282 402 bp DNA linear PAT 26-SEP-2002  
DEFINITION Sequence 176 from patent US 6436387.  
ACCESSION AR223282  
VERSION AR223282.1 GI:23331290  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 402)  
AUTHORS Bauer, S. C., Abrams, M. A., Braford-Goldberg, S. R., Caparon, M. H., Easton, A. M., Klein, B. K., McKearn, J. P., Oling, P. O., Paik, K. and Thomas, J. W.  
TITLE Methods of ex-vivo expansion of hematopoietic cells using multivariant IL-3 hematopoiesis chimera proteins  
JOURNAL Patent: US 6436387-A 176 20-AUG-2002;  
FEATURES Location/Qualifiers  
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Pred. No.: 1.12e-59 Length: 402  
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DB: 6 Gaps: 0  
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Qy 35 GlnGluAlaArgGluLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
Db 61 CAGGAGCCCGGGCTCTCTGAACTGAGTAGACACTGCTGCTGAGATGAATGAACA 120  
Qy 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
Db 121 GTAGAAGTGTATCAGAAATGTTGACCTCCAGAGCCGACTTGCCTACAGACCCGCTG 180  
Qy 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 181 GAGCTGTACAAAGCAGGCGCTTCCCGGCGAGCCTCACCAGCTCAAGGCGCCCTTGACCATG 240  
Qy 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
Db 241 ATGGCCAGCCACTTACAAAGCAGCACTGCCCTCCAAACCCCGGAAACTTCTCTGTGCAACCCAG 300  
Qy 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPhe 134  
Db 301 ATATCACCTTTGAAAGTTTCAAGAGAACCTGGAAGACTTCTGCTGTGTCATCCCTTT 360  
Qy 135 AspCysTrpGluProValGlnGlu 142  
Db 361 GACTGCTGGAGCCAGTCCAGGAG 384  
RESULT 6  
LOCUS A00367 415 bp DNA linear PAT 09-MAR-1993

DEFINITION Artificial gene for granulocyte/macrophage colony stimulating factor.

ACCESSION A00367

VERSION A00367.1 GI:344179

KEYWORDS granulocyte-macrophage colony stimulating factor; synthetic gene.

SOURCE synthetic construct

ORGANISM synthetic construct

REFERENCE 1 (bases 1 to 415)

AUTHORS Synthetic-gene

TITLE Patent: GB 2212159-A 2 19-JUL-1989;

JOURNAL Location/Qualifiers

FEATURES source

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CDS

14..400

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ORIGIN

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Percent Similarity: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0

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QY 35 GlnGluAlaArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54

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QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94

DB 194 GAGCTGTACAGCAGCGCCCTGCGGCGCAGCTCACCAGCTCAGGCGCCCTTGACCATG 253

QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114

DB 254 ATGGCCAGCCACTACAAAGCAGCACTGCCCTCCAAACCCCGGAAACTTCTGTGCAACCCAG 313

QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134

DB 314 ATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTCTGCTGTGTCATCCCTTT 373

QY 135 AspCysTrpGluProValGlnGlu 142

DB 374 GACTGCTGGAGCCAGTCCAGGAG 397

RESULT 7

A00368/c

LOCUS Artificial gene for granulocyte/macrophage colony stimulating factor, reverse complement.

DEFINITION

ACCESSION A00368

VERSION A00368.1 GI:344181

KEYWORDS synthetic construct

SOURCE synthetic construct

ORGANISM synthetic construct

REFERENCE 1 (bases 1 to 415)

AUTHORS Synthetic-gene

TITLE Patent: GB 2212159-A 3 19-JUL-1989;

JOURNAL Location/Qualifiers

FEATURES source

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ORIGIN

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Pred. No.: 1.16e-59 Length: 415

Score: 678.00 Matches: 128

Percent Similarity: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0

Query Match: 88.63% Indels: 0

DB: 6 Gaps: 0

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QY 15 MetAlaProAlaArgSerProSerThrGlnProThrGluHisValAsnAlaIle 34

DB 402 ATGGCACCCTGGGCGCTACCCAGCCAGCAGCCCTGGGAGCATGTGATGCCATC 343

QY 35 GlnGluAlaArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54

DB 342 CAGGAGCCCGCGGTCTCTCTGAACCTGAGTAGACACTGCTGCTGAGATGAATGAACA 283

QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74

DB 282 GTAGAAGTGATATCAGAAATGTTTGACCTCCAGGAGCCGACTTGCCTACAGACCCGCGCTG 223

QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94

DB 222 GAGCTGTACAGCAGCGCCCTGCGGCGCAGCTCACCAGCTCAGGCGCCCTTGACCATG 163

QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114

DB 162 ATGGCCAGCCACTACAAAGCAGCACTGCCCTCCAAACCCCGGAAACTTCTGTGCAACCCAG 103

QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134

DB 102 ATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTCTGCTGTGTCATCCCTTT 43

QY 135 AspCysTrpGluProValGlnGlu 142

DB 42 GACTGCTGGAGCCAGTCCAGGAG 19

RESULT 8

A14305

LOCUS GM-CSF gene (from Homo sapiens).

DEFINITION

ACCESSION A14305

VERSION A14305.1 GI:491775

KEYWORDS synthetic construct

ORGANISM synthetic construct

REFERENCE 1 (bases 1 to 415)

AUTHORS DNA sequence

TITLE Patent: GB 2212160-A 2 19-JUL-1989;

JOURNAL Location/Qualifiers

FEATURES source

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QY      55  ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74
      Db      |||
      559  GTAGAAGTGATATCAGAAATGTTTGACCTCCAGAGCGGACTTGCCCTACAGACCGCGCTG 618
QY      75  GluLeuTyrlsGlnGlySerLeuThrLysLeuLysGlyProLeuThrMet 94
      Db      |||
      619  GAGCTGTACAGCAGCGGCTGCGGGCAGCGCTCACCAGCTCAAGGGCCCCCTTGACCATG 678
QY      95  MetAlaSerHisTyrlsGlnHisCysProThrProGluThrSerCysAlaThrGln 114
      Db      |||
      679  ATGGCCAGCCACTACAGCAGCACTGCCCTCCACCCCGGAACTTCTCTGTGCAACCCAG 738
QY      115  IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134
      Db      |||
      739  ATTATCACCTTTGAAAGTTTCAAAGAGAACCTGAGGACTTCTGCTTGCTCATCCCTTT 798
QY      135  AspCysTrpGluProValGlnGlu 142
      Db      |||
      799  GACTGCTGGAGCCAGTCCAGGAG 822

RESULT 11
AR223222
LOCUS      AR223222      822 bp      DNA      linear      PAT 26-SEP-2002
DEFINITION Sequence 69 from patent US 6436387.
ACCESSION AR223222
VERSION   AR223222.1 GI:23331230
KEYWORDS
SOURCE    Unknown.
ORGANISM  Unclassified.
REFERENCE 1 (bases 1 to 822)
AUTHORS   Bauer, S.C., Abrams, M.A., Braford-Goldberg, S.R., Caparon, M.H.,
          Easton, A.M., Klein, B.K., McKearn, J.P., Olins, P.O., Paik, K. and
          Thomas, J.W.
TITLE     Methods of ex-vivo expansion of hematopoietic cells using
          multivariant IL-3 hematopoiesis chimera proteins
JOURNAL   Patent: US 6436387-A 69 20-AUG-2002;
FEATURES   Location/Qualifiers
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DB:             6      Gaps:         0

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QY      55  ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74
      Db      |||
      559  GTAGAAGTGATATCAGAAATGTTTGACCTCCAGGAGCGGACTTGCCCTACAGACCGCGCTG 618
QY      75  GluLeuTyrlsGlnGlySerLeuThrLysLeuLysGlyProLeuThrMet 94
      Db      |||
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QY      95  MetAlaSerHisTyrlsGlnHisCysProThrProGluThrSerCysAlaThrGln 114
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QY      115  IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134
      Db      |||
      739  ATTATCACCTTTGAAAGTTTCAAAGAGAACCTGAGGACTTCTGCTTGCTCATCCCTTT 798
QY      135  AspCysTrpGluProValGlnGlu 142
      Db      |||
      799  GACTGCTGGAGCCAGTCCAGGAG 822

RESULT 12
AR202217
LOCUS      AR202217      903 bp      DNA      linear      PAT 20-APR-2002
DEFINITION Sequence 66 from patent US 6361977.
ACCESSION AR202217
VERSION   AR202217.1 GI:20256756
KEYWORDS
SOURCE    Unknown.
ORGANISM  Unclassified.
REFERENCE 1 (bases 1 to 903)
AUTHORS   Bauer, S.Christopher., Abrams, M.Allen., Braford-Goldberg, S.Ruth.,
          Caparon, M.Helena., Easton, A.Michael., Klein, B.Kure., McKearn, J.P.,
          Olins, P.O., Paik, K. and Thomas, J.W.
TITLE     Methods of using multivariant IL-3 hematopoiesis fusion protein
JOURNAL   Patent: US 6361977-A 66 26-MAR-2002;
FEATURES   Location/Qualifiers
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      Db      |||
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QY      55  ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74
      Db      |||
      640  GTAGAAGTGATATCAGAAATGTTTGACCTCCAGGAGCGGACTTGCCCTACAGACCGCGCTG 699
QY      75  GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94
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DEFINITION Sequence 66 from patent US 6436387.
ACCESSION AR223219
VERSION AR223219.1 GI:23331227
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 903)
AUTHORS Bauer, S.C., Abrams, M.A., Braford-Goldberg, S.R., Caparon, M.H.,
Easton, A.M., Klein, B.K., McKearn, J.P., Olin, P.O., Paik, K. and
Thomas, J.W.
TITLE Methods of ex-vivo expansion of hematopoietic cells using
multivariant IL-3 hematopoiesis chimera proteins
JOURNAL Patent: US 6436387-A 66 20-AUG-2002;
FEATURES
source
1..903
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QY 35 GlnGluAlaArgArgLeuLeuAenLeuSerArgAspThrAlaAlaGluMetAenGluThr 54
Db 580 CAGGAGCGCGGGGTCTCTCTGAACCTGAGTAGAGACACTGCTCTGAGATGAATGAACA 639
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74
Db 640 GTAGAAGTGAATACAGAAATGTTTGACCTCCAGAGCGGACTTGCCTACAGCCGCCCTG 699
QY 75 GluLeuThrlyrsGlnGlyLeuArgGlySerLeuThrlyrsLeuLyysGlyProLeuThrMet 94
Db 700 GAGCTGTACAGCAGGCGCTCGGGGCGAGCTCACCAGCTCAAGGGCCCTTGACCATG 759
QY 95 MetAlaSerHisTyrlyrsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114
Db 760 ATGGCCAGCCACTACAAGCAGCAGCTGCGCTCCAAACCCCGGAAACTTCCTGTGCAACCCAG 819
QY 115 IleIleThrPheGluSerPheLyysGluAenLeuLyysAspPheLeuLeuValIleProPhe 134
Db 820 ATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTCCTGTGTGTGTCATCCCTTT 879
QY 135 AspCysTrpGluProValGlnGlu 142
Db 880 GACTGTGGGAGCCAGTCCAGGAG 903
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DEFINITION Heterominitobodies.
ACCESSION BD222938
VERSION BD222938.1 GI:33032708
KEYWORDS JP 2002521053-A/32.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 1630)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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AUTHORS Kufer, P., Dreier, T., Baeuerle, P.A., Borschert, K. and Zettl, F.
TITLE Heterominitobodies
PATENT: JP 2002521053-A 32 16-JUL-2002;
MICROMET AG
COMMENT
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PN Mus musculus (mouse)
JP 2002521053-A/32
PD 16-JUL-2002
PF 28-JUL-1999 JP 2000562401
PR 28-JUL-1998 EP 98114082.5
PI PETER KUFER, TORSTEN DREIER, PATRICK A BAEUERLE, KATRIN
BORSCHERT.
PI FLORIAN ZETTL
PC C12N15/09, A61K35/76, A61K38/00, A61K38/21, A61P35/00, A61P35/02,
PC C07K19/00,
PC C12N5/10, C12P21/02, G01N33/53, G01N33/53// (C12N5/10, C12R1:91),
PC (C12P21/02, C12R1:91), C12N15/00, C12N5/00, A61K37/02, A61K37/66,
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CC Heterominitobodies
FH Key Location/Qualifiers
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Db 1260 GCATCCAGAGGCGCGCGCTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAAT 1319
QY 53 GluThrValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThr 72
Db 1320 GAAACAGTAGAGTCACTCAGAAATGTTTGACCTCCAGAGCGGACCTGCCTACAGACC 1379
QY 73 ArgLeuGluLeuThrlyrsGlnGlyLeuArgGlySerLeuThrlyrsLeuLyysGlyProLeu 92
Db 1380 CGCCTGGAGCTGTACAAGCAGGCGCTCGGGGCGAGCTCACCAGCTCAAGGGCCCTTG 1439
QY 93 ThrMetMetAlaSerHisTyrlyrsGlnHisCysProProThrProGluThrSerCysAla 112
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QY 113 ThrGlnIleIleThrPheGluSerPheLyysGluAenLeuLyysAspPheLeuLeuValIle 132
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QY 133 ProPheAspCysTrpGluProValGlnGlu 142
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DEFINITION Sequence 36 from Patent WO0006605.
ACCESSION AX023365
VERSION AX023365.1 GI:10183777
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
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REFERENCE Kufer, P., Zettl, F., Dreier, T., Baeuerle, P.A. and Borschert, K.  
AUTHORS Heteromimibodies  
TITLE Patent: WO 006605-A 36 10-FEB-2000;  
JOURNAL KUFER PETER (DE); ZETTL FLORIAN (DE); DREIER TORSTEN (DE);  
BAEUEERLE PATRICK A (DE); BORSCHERT KATRIN (DE); MICROMET GES FUER  
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## ORIGIN

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QY 33 AlaIleGlnLuhAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsn 52  
DB 1260 GCCATCCAGGAGCGCGGCTCTCTGAACCTGAGTAGACACTGCTGCTGAGATGAAT 1319  
QY 53 GluThrValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThr 72  
DB 1320 GAAACAGTAGAGTCACTCAGAAATGTTTGACCTCCAGAGCCGACCTGCCTACAGACC 1379  
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DB 1380 CGCCTGGAGCTGTACAAGCAGGCGCTCGGGGAGGCTCACCAAGCTCAAGGGCCCTTG 1439  
QY 93 ThrMetMetAlaSerHisTyrllysGlnHisCysProProThrProGluThrSerCysAla 112  
DB 1440 ACCATGATGGCAGGCACCTACAAGCAGACACTGCCCTCCAAACCCCGGAAACTTCCTGTGCA 1499  
QY 113 ThrGlnIleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIle 132  
DB 1500 ACCGATTTATCACCTTTGAAAGTTTCAAGAGAACCTGAGGACTTCTGCTGTTCATC 1559  
QY 133 ProPheAspCysTrpGluProValGlnGlu 142  
DB 1560 CCCTTTGACTGCTGGGAGCCAGTCCAGGAG 1589

Search completed: March 11, 2005, 19:37:25



GenCore version 5.1.6  
Copyright (c) 1993 - 2005 Compugen Ltd.

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Run on: March 11, 2005, 18:21:21 ; Search time 163 Seconds  
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Searched: 1202784 seqs, 818138359 residues

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Post-processing: Minimum Match 0%  
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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	682	89.2	777	3	US-08-762-227A-55
5	682	89.2	777	5	PCT-US95-01185-55
6	678	88.6	402	3	US-08-469-318-176
7	678	88.6	402	3	US-08-468-609A-176
8	678	88.6	402	3	US-08-468-872A-176
9	678	88.6	402	3	US-08-762-227A-176
10	678	88.6	402	5	PCT-US95-01185-176
11	678	88.6	822	3	US-08-469-318-69
12	678	88.6	822	3	US-08-468-609A-69

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14	678	88.6	822	3	US-08-762-227A-69	Sequence 69, Appl
15	678	88.6	822	5	PCT-US95-01185-69	Sequence 69, Appl
16	678	88.6	903	3	US-08-469-318-66	Sequence 66, Appl
17	678	88.6	903	3	US-08-468-609A-66	Sequence 66, Appl
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19	678	88.6	903	3	US-08-762-227A-66	Sequence 66, Appl
20	678	88.6	903	5	PCT-US95-01185-66	Sequence 66, Appl
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22	673	88.0	435	3	US-08-848-760B-8	Sequence 8, Appl
23	673	88.0	435	4	US-09-826-025-8	Sequence 8, Appl
24	673	88.0	435	4	US-10-188-056-31	Sequence 31, Appl
25	673	88.0	435	4	US-10-188-056-32	Sequence 32, Appl
26	673	88.0	660	6	5391485-2	Patent No. 5391485
27	673	88.0	660	6	5391485-2	Patent No. 5391485
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32	673	88.0	900	1	US-08-318-193-7	Sequence 7, Appl
33	673	88.0	905	6	5200327-3	Patent No. 5200327
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35	673	88.0	909	1	US-08-318-193-9	Sequence 9, Appl
36	673	88.0	1011	4	US-09-976-594-275	Sequence 275, App
37	673	88.0	1318	3	US-09-310-842-3	Sequence 3, Appl
38	673	88.0	1588	2	US-09-146-283-1	Sequence 1, Appl
39	673	88.0	1588	3	US-08-579-823A-1	Sequence 1, Appl
40	673	88.0	1588	3	US-09-344-195-1	Sequence 1, Appl
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42	673	88.0	2385	3	US-08-579-823A-3	Sequence 3, Appl
43	673	88.0	2385	3	US-09-344-195-3	Sequence 3, Appl
44	669	87.5	495	6	5405952-1	Patent No. 5405952
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ALIGNMENTS

RESULT 1

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; GENERAL INFORMATION:  
; APPLICANT:  
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion  
; TITLE OF INVENTION: Protein  
; NUMBER OF SEQUENCES: 196  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.30 (EPO)  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/469,318  
; FILING DATE:  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/446,872  
; FILING DATE:  
; INFORMATION FOR SEQ ID NO: 55:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 777 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
US-08-469-318-55  
  
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Query Match: 89.15% Indels: 0

DB: 3 Gaps: 0  
US-10-723-083-2 (1-142) x US-08-469-318-55 (1-777)  
QY 9 SerGlyLeuGluArgMetAlaProAlaArgSerProSerProSerThrGlnProTrp 28  
Db 376 TCTGGCGGCGCTCCAACTGGCAGCGGCTCGTTCCCGCTCCCGCTACCCAGCGGTGG 435  
QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAla 48  
Db 436 GAACACGTGAATGCCATCCAGGAGCGCGGCTCTCTGAACCTGAGTAGAGACTGCT 495  
QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnGluProThr 68  
Db 496 GCTGAGATGAATGAACAGTAGAGTATCAGAAATGTTTGACCTCCAGGAGCGGACT 555  
QY 69 CysLeuGlnThrArgLeuGluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeu 88  
Db 556 TGCCTACAGACCGCGCTGGAGCTGTACAAGCAGGGCGCTGCGGGGCGAGCTCACCAAGCTC 615  
QY 89 LysGlyProLeuThrMetAlaSerHisTyrlsGlnHisCysProProThrProGlu 108  
Db 616 AAGGGCCCTTGACCATGATGGCCAGCCACTACAGCAGACTGCGCTCCAAACCCCGGAA 675  
QY 109 ThrSerCysAlaThrGlnIleIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128  
Db 676 ACTTCTGTGTCAACCCAGATTATCACCTTTGAAAGTTTCAAAGAGAACTTGAGGACTTC 735  
QY 129 LeuLeuValIleProPheAspCysTrpGluProValGlnGlu 142  
Db 736 CTGCTGTATCCCTTTGACTGCTGGAGCCAGTCCAGGAG 777

## RESULT 2

US-08-468-609A-55  
; Sequence 55, Application US/08468609A  
; Patent No. 6030812  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; APPLICANT: Bauer, S. C.  
; APPLICANT: Braford-Goldberg, Sarah R.  
; APPLICANT: Caparon, Mair H.  
; APPLICANT: Easton, Alan M.  
; APPLICANT: Klein, Barbara K.  
; APPLICANT: McKearn, John P.  
; APPLICANT: Olin, Peter O.  
; APPLICANT: Paik, Kuman  
; APPLICANT: Thomas, John W.  
; TITLE OF INVENTION: Fusion Proteins Comprising Multiply Mutated Inteleukin-3 (II-  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
; ADDRESSEE: Corporate Patent Dept.  
; STREET: P. O. Box 5110  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60680  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/468,609A  
; FILING DATE: 06-JUN-1995  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Bennett, Dennis A.  
; REGISTRATION NUMBER: 34,547  
; REFERENCE/DOCKET NUMBER: C-2790/3

## TELECOMMUNICATION INFORMATION:

; TELEPHONE: (314)737-6986  
; TELEFAX: (314)737-6972  
; INFORMATION FOR SEQ ID NO: 55:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 777 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; US-08-468-609A-55

## Alignment Scores:

Pred. No.: 1-89e-76 Length: 777  
Score: 682.00 Matches: 130  
Percent Similarity: 97.01% Conservative: 0  
Best Local Similarity: 97.01% Mismatches: 4  
Query Match: 89.15% Indels: 0  
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-468-609A-55 (1-777)

QY 9 SerGlyLeuGluArgMetAlaProAlaArgSerProSerProSerThrGlnProTrp 28  
Db 376 TCTGGCGGCGCTCCAACTGGCAGCGGCTCGTTCCCGCTCCCGCTACCCAGCGGTGG 435  
QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAla 48  
Db 436 GAACACGTGAATGCCATCCAGGAGCGCGGCTCTCTGAACCTGAGTAGAGACTGCT 495  
QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnGluProThr 68  
Db 496 GCTGAGATGAATGAACAGTAGAGTATCAGAAATGTTTGACCTCCAGGAGCGGACT 555  
QY 69 CysLeuGlnThrArgLeuGluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeu 88  
Db 556 TGCCTACAGACCGCGCTGGAGCTGTACAAGCAGGGCGCTGCGGGGCGAGCTCACCAAGCTC 615  
QY 89 LysGlyProLeuThrMetAlaSerHisTyrlsGlnHisCysProProThrProGlu 108  
Db 616 AAGGGCCCTTGACCATGATGGCCAGCCACTACAGCAGACTGCGCTCCAAACCCCGGAA 675  
QY 109 ThrSerCysAlaThrGlnIleIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128  
Db 676 ACTTCTGTGTCAACCCAGATTATCACCTTTGAAAGTTTCAAAGAGAACTTGAGGACTTC 735  
QY 129 LeuLeuValIleProPheAspCysTrpGluProValGlnGlu 142  
Db 736 CTGCTGTATCCCTTTGACTGCTGGAGCCAGTCCAGGAG 777

## RESULT 3

US-08-446-872A-55  
; Sequence 55, Application US/08446872A  
; Patent No. 6361977  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; APPLICANT: Bauer, S. C.  
; APPLICANT: Braford-Goldberg, Sarah R.  
; APPLICANT: Caparon, Mair H.  
; APPLICANT: Easton, Alan M.  
; APPLICANT: Klein, Barbara K.  
; APPLICANT: McKearn, John P.  
; APPLICANT: Olin, Peter O.  
; APPLICANT: Paik, Kuman  
; APPLICANT: Thomas, John W.  
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis  
; TITLE OF INVENTION: Fusion Protein  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
; ADDRESSEE: Corporate Patent Dept.  
; STREET: P. O. Box 5110  
; CITY: Chicago

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; STATE: Illinois
; COUNTRY: USA
; ZIP: 60680
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; FILING DATE: 06-JUN-1995
; PRIORITY APPLICATION NUMBER: US/08/446,872A
; FILING DATE: 06-JUN-1995
; PRIORITY APPLICATION NUMBER: US/08/192,325
; FILING DATE: 14-FEB-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Bennett, Dennis A.
; REGISTRATION/DOCKET NUMBER: C-2790/1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)737-6986
; TELEFAX: (314)737-6972
; INFORMATION FOR SEQ ID NO: 55:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 777 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-446-872A-55

Alignment Scores:
Pred. No.: 1,89e-76 Length: 777
Score: 682.00 Matches: 130
Percent Similarity: 97.01% Conservative: 0
Best Local Similarity: 97.01% Mismatches: 4
Query Match: 89.15% Indels: 0
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-446-872A-55 (1-777)
QY 9 SerGlyIleGluGlyArgMetAlaProAlaArgSerProSerProSerThrGlnProTyr 28
Db 376 TCTGGCGGGCGGCTCCAACTGGCAGCGGCTCGTTCGCCGCTCCCGCTACCCAGCGGTG 435
QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAla 48
Db 436 GAACACGTGAATGCCATCCAGAGGCGCGGCTCTCTGAACCTGAGTAGAGACACTGCT 495
QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnProThr 68
Db 496 GCTGAGATGAATGAACAGTAGAAGTGATATCAGAAATGTTGACCTCCAGGAGCGGACT 555
QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88
Db 556 TGCCCTACAGACCCGCTGGAGCTGTACAAGCAGGGCTCGGGGGAGCCCTCACAGCTC 615
QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProProThrProGlu 108
Db 616 AAGGGCCCTTGACCATGATGGCCGACCTACAGCAGCACTGCGCTCCAAACCCCGGAA 675
QY 109 ThrSerCysAlaThrGlnIleIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128
Db 676 ACTTCTGTGCAACCCAGATTATCAGCTTGAAGTTTCAAGAGAACTTGAAGGACTTC 735
QY 129 LeuLeuValIleProPheAspCysTyrGluProValGlnGlu 142
Db 736 CTGCTTGTATCCCTTTGACTCTGGAGCCAGTCCAGGAG 777

RESULT 4
US-08-762-227A-55
; Sequence 55, Application US/08762227A
; Patent No. 6436387

; GENERAL INFORMATION:
; APPLICANT: Abrams, Mark A.
; Bauer, S. C.
; Braford-Goldberg, Sarah R.
; Caparon, Maire H.
; Easton, Alan M.
; Klein, Barbara K.
; McKeam, John P.
; Olin, Peter O.
; Paik, Kumnan
; Thomas, John W.
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis
; NUMBER OF SEQUENCES: 197
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,
; Corporate Patent Dept.
; STREET: P. O. Box 5110
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60680
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; FILING DATE: 09-Dec-1996
; APPLICATION NUMBER: US/08/762,227A
; CLASSIFICATION: <Unknown>
;
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/192,325
; FILING DATE: 14-FEB-1994
; APPLICATION NUMBER: US 08/446,872
; FILING DATE: 06-JUN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Bennett, Dennis A.
; REGISTRATION NUMBER: 34,547
; REFERENCE/DOCKET NUMBER: C-2790/5
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (708)470-6501
; TELEFAX: (708)470-8881
; INFORMATION FOR SEQ ID NO: 55:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 777 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; SEQUENCE DESCRIPTION: SEQ ID NO: 55:
; US-08-762-227A-55

Alignment Scores:
Pred. No.: 1,89e-76 Length: 777
Score: 682.00 Matches: 130
Percent Similarity: 97.01% Conservative: 0
Best Local Similarity: 97.01% Mismatches: 4
Query Match: 89.15% Indels: 0
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-762-227A-55 (1-777)
QY 9 SerGlyIleGluGlyArgMetAlaProAlaArgSerProSerProSerThrGlnProTyr 28
Db 376 TCTGGCGGGCGGCTCCAACTGGCAGCGGCTCGTTCGCCGCTCCCGCTACCCAGCGGTG 435
QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAla 48
Db 436 GAACACGTGAATGCCATCCAGAGGCGCGGCTCTCTGAACCTGAGTAGAGACACTGCT 495
QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnProThr 68
Db 496 GCTGAGATGAATGAACAGTAGAAGTGATATCAGAAATGTTGACCTCCAGGAGCGGACT 555
QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88
Db 556 TGCCCTACAGACCCGCTGGAGCTGTACAAGCAGGGCTCGGGGGAGCCCTCACAGCTC 615
QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProProThrProGlu 108
Db 616 AAGGGCCCTTGACCATGATGGCCGACCTACAGCAGCACTGCGCTCCAAACCCCGGAA 675
QY 109 ThrSerCysAlaThrGlnIleIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128
Db 676 ACTTCTGTGCAACCCAGATTATCAGCTTGAAGTTTCAAGAGAACTTGAAGGACTTC 735
QY 129 LeuLeuValIleProPheAspCysTyrGluProValGlnGlu 142
Db 736 CTGCTTGTATCCCTTTGACTCTGGAGCCAGTCCAGGAG 777

RESULT 4
US-08-762-227A-55
; Sequence 55, Application US/08762227A
; Patent No. 6436387
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Db 496 GCTGAGATGAATGAACAGTAGAGTAGATATCAGAAATGTTTGACCTCCAGGAGCCGACT 555  
QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88  
Db 556 TGCTACAGACCCGCTGGAGCTGTACACAGAGGCTCGGGGCGAGCCTCACCAGGCTC 615  
QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProThrProGlu 108  
Db 616 AAGGGCCCTTGACATGATGGCCAGCCACTACAAGCAGCACTGCCCTCCAAACCCGGAA 675  
QY 109 ThrSerCysAlaThrGlnIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128  
Db 676 ACTTCCTGTGCAACCCAGATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTC 735  
QY 129 LeuLeuValIleProPheAspCysTrpGluProValGlnGlu 142  
Db 736 CTGCTGTGATCCCTTTGACTGCTGGAGCCAGTCAGGAG 777

## RESULT 5

PCT-US95-01185-55  
; Sequence 55, Application PC/TUS9501185  
; GENERAL INFORMATION:  
; APPLICANT:  
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion  
; TITLE OF INVENTION: Protein  
; NUMBER OF SEQUENCES: 196  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.30 (EPO)  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: PCT/US95/01185  
; FILING DATE: 02-FEB-1995  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192325  
; FILING DATE: 14-FEB-1994  
; INFORMATION FOR SEQ ID NO: 55:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 777 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
PCT-US95-01185-55

Alignment Scores:  
Pred. No.: 1.89e-76 Length: 777  
Score: 682.00 Matches: 130  
Percent Similarity: 97.01% Conservative: 0  
Best Local Similarity: 97.01% Mismatches: 4  
Query Match: 89.15% Indels: 0  
DB: 5 Gaps: 0

US-10-723-083-2 (1-142) x PCT-US95-01185-55 (1-777)

QY 9 SerGlyIleGluGlyArgMetAlaProAlaArgSerProSerThrGlnProTrp 28  
Db 376 TCTGGCGGCGCTCCAACTGGCACCGCTCGTTCCCGCTCCCGCTTACCCAGCGCTGG 435  
QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAla 48  
Db 436 GAACAGTGAATGCCATCCAGAGGCGCGCGCTCTCTGAACCTGAGTAGACACTGCT 495  
QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnProThr 68  
Db 496 GCTGAGATGAATGAACAGTAGAGTAGATATCAGAAATGTTTGACCTCCAGGAGCCGACT 555  
QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88  
Db 556 TGCTACAGACCCGCTGGAGCTGTACAGAGGCGCTCGGGGCGAGCCTCACCAGGCTC 615

QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProThrProGlu 108  
Db 616 AAGGGCCCTTGACATGATGGCCAGCCACTACAAGCAGCACTGCCCTCCAAACCCGGAA 675  
QY 109 ThrSerCysAlaThrGlnIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128  
Db 676 ACTTCCTGTGCAACCCAGATTATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTC 735  
QY 129 LeuLeuValIleProPheAspCysTrpGluProValGlnGlu 142  
Db 736 CTGCTGTGATCCCTTTGACTGCTGGAGCCAGTCAGGAG 777

## RESULT 6

US-08-469-318-176  
; Sequence 176, Application US/08469318  
; Patent No. 6022535  
; GENERAL INFORMATION:  
; APPLICANT:  
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion  
; TITLE OF INVENTION: Protein  
; NUMBER OF SEQUENCES: 196  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.30 (EPO)  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/469,318  
; FILING DATE:  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/446,872  
; FILING DATE:  
; INFORMATION FOR SEQ ID NO: 176:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 402 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
US-08-469-318-176

Alignment Scores:  
Pred. No.: 2.23e-76 Length: 402  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-469-318-176 (1-402)

QY 15 MetAlaProAlaArgSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
Db 1 ATGGCACCGGCTCGTTCCCGCTCCCGCTTACCCAGCGCTGGGAAACAGTGAATCCCATC 60  
QY 35 GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
Db 61 CAGGAGGCGCGGCTCTCTGAACTGAGTAGAGACACTGCTGCTGAGATGAATGAACA 120  
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
Db 121 GTAGAAGTGATATCAGAAATGTTTGACCTCCAGGAGCGGACTTGCTTACAGACCCGCTG 180  
QY 75 GluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 181 GAGCTGTAAACAGAGCGGCTCGGGGCGAGCTTCAACAGCTCAAGGGCCCTTGGACCATG 240  
QY 95 MetAlaSerHisTyrLysGlnHisCysProThrProGluThrSerCysAlaThrGln 114  
Db 241 ATGGCAGCCACTACAAGCAGCACTGCCCTCCAAACCCGGGAAACTTCTGTGTCAACCCAG 300  
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134

Db 301 ATTATCACCTTTGAAAGTTTCAAGAGAACTCGAGGACTTCTGCTGTGTCATCCCTTT 360  
Qy 135 AspCysTrpGluProValGlnGlu 142  
Db 361 GACTGCTGGGAGCCAGTCCAGGAG 384

RESULT 7  
US-08-468-609A-176  
; Sequence 176, Application US/08468609A  
; Patent No. 6030812  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; APPLICANT: Bauer, S. C.  
; APPLICANT: Braford-Goldberg, Sarah R.  
; APPLICANT: Caparon, Mair H.  
; APPLICANT: Easton, Alan M.  
; APPLICANT: Klein, Barbara K.  
; APPLICANT: McKearn, John P.  
; APPLICANT: Olin, Peter O.  
; APPLICANT: Olin, Peter O.  
; APPLICANT: Thomas, John W.  
; TITLE OF INVENTION: Fusion Proteins Comprising Multiply Mutated Inteleukin-3 (IL-  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
; ADDRESSEE: Corporate Patent Dept.  
; STREET: P. O. Box 5110  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60680

; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; FILING DATE: 06-JUN-1995  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Bennett, Dennis A.  
; REGISTRATION NUMBER: 34,547  
; REFERENCE/DOCKET NUMBER: C-2790/3  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (314)737-6986  
; TELEFAX: (314)737-6972  
; INFORMATION FOR SEQ ID NO: 176:  
; SEQUENCE CHARACTERISTICS:  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
US-08-468-609A-176

Alignment Scores:  
Pred. No.: 2,23e-76 Length: 402  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-468-609A-176 (1-402)  
Qy 15 MetAlaProAlaArgSerProSerThrGlnProThrGluHisValAsnAlaIle 34  
Db 1 ATGGCACCGGCTGTTCCCGTCTACCCAGCCGTTGGGAACACGATGCAATGCCATC 60

Qy 35 GlnGluAlaArgSerArgSerLeuSerArgSerThrAlaAlaGluMetAsnGluThr 54  
Db 61 CAGGAGGCCCGCGTCTCTGAACTGAGTAGAGACACTGCTGCTGAGATGAACA 120  
Qy 55 ValGluValIleSerGluMetPheAspLeuGlnProThrCysLeuGlnThrArgLeu 74  
Db 121 GTAGAAAGTGTATCAGAAATGTTTGACCTCCAGGAGCCGACTTGCCTACAGACCCGCTG 180  
Qy 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 181 GAGTGTACAAAGCAGGCGCTGCGGGGCGCCCTCACCAGCTCAAGGGCCCTTGACCATG 240  
Qy 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
Db 241 ATGGCCAGCCACTACAAAGCAGCAGCTGCCCTCCAAACCCGGAACACTTCTGTGCAACCCAG 300  
Qy 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
Db 301 ATTATCACCTTTGAAAGTTTCAAGAGAACTTCTGCTGTGTCATCCCTTT 360  
Qy 135 AspCysTrpGluProValGlnGlu 142  
Db 361 GACTGCTGGGAGCCAGTCCAGGAG 384

RESULT 8  
US-08-446-872A-176  
; Sequence 176, Application US/08446872A  
; Patent No. 6361977  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; APPLICANT: Bauer, S. C.  
; APPLICANT: Braford-Goldberg, Sarah R.  
; APPLICANT: Caparon, Mair H.  
; APPLICANT: Easton, Alan M.  
; APPLICANT: Klein, Barbara K.  
; APPLICANT: McKearn, John P.  
; APPLICANT: Olin, Peter O.  
; APPLICANT: Olin, Peter O.  
; APPLICANT: Thomas, John W.  
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis  
; TITLE OF INVENTION: Fusion Protein  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
; ADDRESSEE: Corporate Patent Dept.  
; STREET: P. O. Box 5110  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60680  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; FILING DATE: 06-JUN-1995  
; CLASSIFICATION: 424  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Bennett, Dennis A.  
; REGISTRATION NUMBER: 34,547  
; REFERENCE/DOCKET NUMBER: C-2790/1  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (314)737-6986  
; TELEFAX: (314)737-6972  
; INFORMATION FOR SEQ ID NO: 176:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 402 base pairs

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; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-446-872A-176

Alignment Scores:
Pred. No.: 2,23e-76 Length: 402
Score: 678.00 Matches: 128
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 88.63% Indels: 0
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-446-872A-176 (1-402)
QY 15 MetAlaProAlaArgSerProSerThrClnProTrpGluHisValAsnAlaIle 34
Db 1 ATGGCACCGGCTGTTCCCGTCTACCCAGCGTGGGAACACGTAATGCCATC 60
QY 35 GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54
Db 61 CAGGAGCGCGGCTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAACA 120
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74
Db 121 GTAGAAGTGATATCAGAAATGTTGACCTCCAGGAGCGGACTTGCCTACAGACCGCGCTG 180
QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94
Db 181 GAGCTGTACAGCAGGCGCTGCGGGCAGCGCTACCAAGCTCAAGGCGCCCTTGACCATG 240
QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114
Db 241 ATGGCCAGCCACTACAAGCAGCACTGCCCTCCAAACCCCGGAACTTCTCTGTCGCAACCCAG 300
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134
Db 301 ATTATCACCTTTGAAAGTTTCAAGAGAACCTGAGGACTTCTGCTTGTATCCCTTT 360
QY 135 AspCysTrpGluProValGlnGlu 142
Db 361 GACTGTGGGAGCCAGTCCAGGAG 384

RESULT 9
US-08-762-227A-176
; Sequence 176, Application US/08762227A
; Patent No. 6436387
; GENERAL INFORMATION:
; APPLICANT: Abrams, Mark A.
; Bauer, S. C.
; Braford-Goldberg, Sarah R.
; Caparon, Maïre H.
; Easton, Alan M.
; Klein, Barbara K.
; McKearn, John P.
; Ollins, Peter O.
; Paik, Kumnan
; Thomas, John W.
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis
; Fusion Protein
; NUMBER OF SEQUENCES: 197
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,
; Corporate Patent Dept.
; STREET: P. O. Box 5110
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60680
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA: US/08/762,227A
APPLICATION NUMBER: US/08/762,227A
FILING DATE: 09-Dec-1996
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/192,325
FILING DATE: 14-FEB-1994
APPLICATION NUMBER: US 08/446,872
FILING DATE: 06-JUN-1995
ATTORNEY/AGENT INFORMATION:
NAME: Bennett, Dennis A.
REGISTRATION NUMBER: 34,547
REFERENCE/DOCKET NUMBER: C-2790/5
TELECOMMUNICATION INFORMATION:
TELEPHONE: (708) 470-6501
TELEFAX: (708) 470-6881
INFORMATION FOR SEQ ID NO: 176:
SEQUENCE CHARACTERISTICS:
LENGTH: 402 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
SEQUENCE DESCRIPTION: SEQ ID NO: 176:
US-08-762-227A-176

Alignment Scores:
Pred. No.: 2,23e-76 Length: 402
Score: 678.00 Matches: 128
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 88.63% Indels: 0
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-762-227A-176 (1-402)
QY 15 MetAlaProAlaArgSerProSerThrGlnProTrpGluHisValAsnAlaIle 34
Db 1 ATGGCACCGGCTGTTCCCGTCTACCCAGCGTGGGAACACGTAATGCCATC 60
QY 35 GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54
Db 61 CAGGAGCGCGGCTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAACA 120
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74
Db 121 GTAGAAGTGATATCAGAAATGTTGACCTCCAGGAGCGGACTTGCCTACAGACCGCGCTG 180
QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94
Db 181 GAGCTGTACAGCAGGCGCTGCGGGCAGCGCTACCAAGCTCAAGGCGCCCTTGACCATG 240
QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114
Db 241 ATGGCCAGCCACTACAAGCAGCACTGCCCTCCAAACCCCGGAACTTCTCTGTCGCAACCCAG 300
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134
Db 301 ATTATCACCTTTGAAAGTTTCAAGAGAACCTGAGGACTTCTGCTTGTATCCCTTT 360
QY 135 AspCysTrpGluProValGlnGlu 142
Db 361 GACTGTGGGAGCCAGTCCAGGAG 384

RESULT 10
PCT-US95-01185-176
; Sequence 176, Application PC/TUS9501185
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion
; Protein
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NUMBER OF SEQUENCES: 196  
COMPUTER READABLE FORM: disk  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/01185  
FILING DATE: 02-FEB-1995  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/192325  
FILING DATE: 14-FEB-1994  
INFORMATION FOR SEQ ID NO: 176:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 402 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
PCT-US95-01185-176

Alignment Scores:  
Pred. No.: 2,23e-76 Length: 402  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 5 Gaps: 0

US-10-723-083-2 (1-142) x PCT-US95-01185-176 (1-402)

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QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34
DB 1 ATGGCACCGGCTGTTCCCGTCCCGTCTACCCAGCCGCGGGAACACCGTGAATGCCATC 60
QY 35 GlnGluAlaArgArgLeuLeuLeuLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54
DB 61 CAGGAGGCCCGGCGTCTCCTGAACTGAGTACAGACACTGCTGCTGAGATGAATGAACA 120
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74
DB 121 GTAGAAGTGATATCAGAAATGTTGACCTCCAGAGCCGACTTGCCTCAGACCCGCCCTG 180
QY 75 GluLeuTyTrpLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94
DB 181 GAGCTGTACAGCAGGCGCTCGCGGCGAGCCTCACCAGCTCAAGGCGCCCTTGACCATG 240
QY 95 MetAlaSerHisTyTrpLysGlnHisCysProProThrProGluThrSerCysAlaThrGln 114
DB 241 ATGGCCAGGCACCTACCAAGCAGCAGTGCCTCCAAACCCCGGAAACTTCTCTGTGCAACCCAG 300
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPhe 134
DB 301 ATTATCACCTTTGAAAGTTTCAAGAGAACCTTCAAGGACTTCTGCTTGTGTCATCCCTTT 360
QY 135 AspCysTrpGluProValGlnGlu 142
DB 361 GACTGCTGGGAGCCAGCTCCAGGAG 384
```

RESULT 11

US-08-469-318-69  
Sequence 69, Application US/08469318  
Patent No. 6022535  
GENERAL INFORMATION:  
APPLICANT:  
APPLICANT: Bauer, S. C.  
APPLICANT: Braford-Goldberg, Sarah R.  
APPLICANT: Caparon, Maïre H.  
APPLICANT: Easton, Alan M.  
APPLICANT: Klein, Barbara K.  
APPLICANT: McKearn, John P.  
APPLICANT: Oline, Peter O.  
APPLICANT: Paik, Kumnan  
APPLICANT: Thomas, John W.  
TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion  
NUMBER OF SEQUENCES: 196  
COMPUTER READABLE FORM: disk  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/469,318  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/446,872  
FILING DATE:  
INFORMATION FOR SEQ ID NO: 69:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 822 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-469-318-69

Alignment Scores:  
Pred. No.: 6.6e-76 Length: 822  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-469-318-69 (1-822)

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QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34
DB 439 ATGGCACCGGCTGTTCCCGTCCCGTCTACCCAGCCGCGGGAACACCGTGAATGCCATC 498
QY 35 GlnGluAlaArgArgLeuLeuLeuLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54
DB 499 CAGGAGGCCCGGCGTCTCCTGAACTGAGTACAGACACTGCTGCTGAGATGAATGAACA 558
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74
DB 559 GTAGAAGTGATATCAGAAATGTTGACCTCCAGAGCCGACTTGCCTCAGACCCGCCCTG 618
QY 75 GluLeuTyTrpLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94
DB 619 GAGCTGTACAGCAGGCGCTCGCGGCGAGCCTCACCAGCTCAAGGCGCCCTTGACCATG 678
QY 95 MetAlaSerHisTyTrpLysGlnHisCysProProThrProGluThrSerCysAlaThrGln 114
DB 679 ATGGCCAGGCACCTACCAAGCAGCAGTGCCTCCAAACCCCGGAAACTTCTCTGTGCAACCCAG 738
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPhe 134
DB 739 ATTATCACCTTTGAAAGTTTCAAGAGAACCTTCAAGGACTTCTGCTTGTGTCATCCCTTT 798
QY 135 AspCysTrpGluProValGlnGlu 142
DB 799 GACTGCTGGGAGCCAGCTCCAGGAG 822
```

RESULT 12

US-08-468-609A-69  
Sequence 69, Application US/08468609A  
Patent No. 6030812  
GENERAL INFORMATION:  
APPLICANT:  
APPLICANT: Abrams, Mark A.  
APPLICANT: Bauer, S. C.  
APPLICANT: Braford-Goldberg, Sarah R.  
APPLICANT: Caparon, Maïre H.  
APPLICANT: Easton, Alan M.  
APPLICANT: Klein, Barbara K.  
APPLICANT: McKearn, John P.  
APPLICANT: Oline, Peter O.  
APPLICANT: Paik, Kumnan  
APPLICANT: Thomas, John W.  
TITLE OF INVENTION: Fusion Proteins Comprising Multiply Mutated Inteleukin-3 (IL-3)  
NUMBER OF SEQUENCES: 197  
CORRESPONDENCE ADDRESS:

ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
ADDRESSEE: Corporate Patent Dept.,  
STREET: P. O. Box 5110  
CITY: Chicago  
STATE: Illinois  
COUNTRY: USA  
ZIP: 60680  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/468, 609A  
FILING DATE: 06-JUN-1995  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/192,325  
FILING DATE: 14-FEB-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Bennett, Dennis A.  
REGISTRATION NUMBER: 34,547  
REFERENCE/DOCKET NUMBER: C-2790/3  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (314)737-6986  
TELEFAX: (314)737-6972  
INFORMATION FOR SEQ ID NO: 69:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 822 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-468-609A-69

Alignment Scores:  
Pred. No.: 6.6e-76 Length: 822  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-468-609A-69 (1-822)

QY 15 MetAlaProAlaArgSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
Db 439 ATGGCAGCGGCTGTTCCCGTCCCGCTTACCCAGCGCGTGGAAACACGTGATGCCATC 498  
QY 35 GlnGluAlaArgArgLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
Db 499 CAGGAGGCGCGGCTCTCTGAACTGAGTAGAGACACTGCTGAGATGAATGAACA 558  
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
Db 559 GTAGAAGTATATACAGAAATGTTGACCTCCAGAGCGGACTTGCCTCAGACCGCGCTG 618  
QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 619 GAGCTGTACAAGCAGGCGCTCGCGGCGAGCCTCACCAAGCTCAAGGCGCCCTTGACCATG 678  
QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
Db 679 ATGGCCAGGCACCTACAGAGGAGCAGCACTGCCCTCCAAACCCCGGAAACTTCTGTGCAACCCAG 738  
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPhe 134  
Db 739 ATTATCACCTTTGAAGATTTCAAAGAACCTGAGGACTTCTGCTGTGTCATCCCTTT 798  
QY 135 AspCysTrpGluProValGlnGlu 142  
Db 799 GACTGTGGGAGCCAGTCCAGGAG 822

RESULT 13  
US-08-446-872A-69  
Sequence 69, Application US/08446872A  
Patent No. 6361977  
GENERAL INFORMATION:  
APPLICANT: Abrams, Mark A.  
APPLICANT: Bauer, S. C.  
APPLICANT: Braford-Goldberg, Sarah R.  
APPLICANT: Caparon, Mair H.  
APPLICANT: Easton, Alan M.  
APPLICANT: Klein, Barbara K.  
APPLICANT: McKearn, John P.  
APPLICANT: Olins, Peter O.  
APPLICANT: Paik, Kumnan  
APPLICANT: Thomas, John W.  
TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis  
TITLE OF INVENTION: Fusion Protein  
NUMBER OF SEQUENCES: 197  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
STREET: P. O. Box 5110  
CITY: Chicago  
STATE: Illinois  
COUNTRY: USA  
ZIP: 60680  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/446, 872A  
FILING DATE: 06-JUN-1995  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/192,325  
FILING DATE: 14-FEB-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Bennett, Dennis A.  
REGISTRATION NUMBER: 34,547  
REFERENCE/DOCKET NUMBER: C-2790/1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (314)737-6986  
TELEFAX: (314)737-6972  
INFORMATION FOR SEQ ID NO: 69:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 822 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-446-872A-69

Alignment Scores:  
Pred. No.: 6.6e-76 Length: 822  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 3 Gaps: 0

US-10-723-083-2 (1-142) x US-08-446-872A-69 (1-822)

QY 15 MetAlaProAlaArgSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
Db 439 ATGGCAGCGGCTGTTCCCGTCCCGCTTACCCAGCGCGTGGAAACACGTGATGCCATC 498  
QY 35 GlnGluAlaArgArgLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
Db 499 CAGGAGGCGCGGCTCTCTGAACTGAGTAGAGACACTGCTGCTGAGATGAATGAACA 558  
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74



Db 559 GTAGAGTGTATACAGAAATGTTTGACCTCCAGAGCCGACTTGCCTACAGACCCCGCTG 618  
QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 619 GAGCTGTACAAAGCAGGCGCTGCGGGCAGGCTCACCAAGCTCAAGGCGCCCTTGACCATG 678  
QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
Db 679 ATGGCAGGCACCTACCAAGCAGCAGCTCCCTCCAAACCCCGGAAACTTCTCTGTGCAACCCAG 738  
QY 115 IleileThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
Db 739 ATTATCACCTTTGAAGATTTCAAAGAACCTGAGGACTTCTGCTGTGTCATCCCTTT 798  
QY 135 AspCysTrpGluProValGlnGlu 142  
Db 799 GACTGCTGGGAGCCAGTCCAGGAG 822

RESULT 14  
US-08-762-227A-69  
; Sequence 69, Application US/08762227A  
; Patent No. 6436387  
; GENERAL INFORMATION:  
; APPLICANT: Abrams, Mark A.  
; Bauer, S. C.  
; Bradford-Goldberg, Sarah R.  
; Caparon, Mairé H.  
; Easton, Alan M.  
; Klein, Barbara K.  
; McKearn, John P.  
; Ollins, Peter O.  
; Paik, Kumnan  
; Thomas, John W.  
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis  
; FUSION PROTEIN  
; NUMBER OF SEQUENCES: 197  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Dennis A. Bennett, G.D. Searle & Co.,  
; Corporate Patent Dept.  
; STREET: P. O. Box 5110  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60680  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/762,227A  
; FILING DATE: 09-Dec-1996  
; CLASSIFICATION: <Unknown>  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; APPLICATION NUMBER: US 08/446,872  
; FILING DATE: 06-JUN-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Bennett, Dennis A.  
; REGISTRATION NUMBER: 34,547  
; REFERENCE/DOCKET NUMBER: C-2790/5  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (708)470-6501  
; TELEFAX: (708)470-6881  
; INFORMATION FOR SEQ ID NO: 69:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 822 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)

; SEQUENCE DESCRIPTION: SEQ ID NO: 69:  
US-08-762-227A-69  
Alignment Scores:  
Pred. No.: 6,6e-76 Length: 822  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 3 Gaps: 0  
US-10-723-083-2 (1-142) x US-08-762-227A-69 (1-822)  
QY 15 MetAlaProAlaArgSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
Db 439 ATGGCAGCGGCTGTTCCCGCTACCCAGCCGCGGGAACACGTAATGCCATC 498  
QY 35 GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
Db 499 CAGGAGGCCCGCGCTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAATGAACA 558  
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
Db 559 GTAGAAGTGTATACAGAAATGTTTGACCTCCAGGAGCCGACTTGCCTACAGACCCCGCTG 618  
QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 619 GAGCTGTACAAAGCAGGCGCTGCGGGCAGGCTCACCAAGCTCAAGGCGCCCTTGACCATG 678  
QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
Db 679 ATGGCAGGCACCTACCAAGCAGCAGCTCCCTCCAAACCCCGGAAACTTCTCTGTGCAACCCAG 738  
QY 115 IleileThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
Db 739 ATTATCACCTTTGAAGATTTCAAAGAACCTGAGGACTTCTGCTGTGTCATCCCTTT 798  
QY 135 AspCysTrpGluProValGlnGlu 142  
Db 799 GACTGCTGGGAGCCAGTCCAGGAG 822

RESULT 15  
PCT-US95-01185-69  
; Sequence 69, Application PC/TUS9501185  
; GENERAL INFORMATION:  
; APPLICANT:  
; TITLE OF INVENTION: Multivariant IL-3 Hematopoiesis Fusion  
; TITLE OF INVENTION: Protein  
; NUMBER OF SEQUENCES: 196  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.30 (EPO)  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: PCT/US95/01185  
; FILING DATE: 02-FEB-1995  
; CLASSIFICATION:  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 08/192325  
; FILING DATE: 14-FEB-1994  
; INFORMATION FOR SEQ ID NO: 69:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 822 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; PCT-US95-01185-69  
Alignment Scores:  
Pred. No.: 6,6e-76 Length: 822  
Score: 678.00 Matches: 128

Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	88.63%	Indels:	0
DB:	5	Gaps:	0

US-10-723-083-2 (1-142) x PCT-US95-01185-69 (1-822)

Qy	15	MetAlaProAlaArgSerProSerProSerThrGlnProThrGluHisValAsnAlaIle	34
Db	439	ATGGCACCGCGCTCGTTCCCGCTCCCGCTACCCAGCGCTGGGAACACGTGAATGCATC	498
Qy	35	GlnGluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr	54
Db	499	CAGGAGCGCGCGGCTCTCTGAACTGAGTAGACACACTGCTGCTGAGATGAATGAACA	558
Qy	55	ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu	74
Db	559	GTAGAAGTCAATATCAGAAATGTTTGACCTCCAGGAGCGGACTGGCTTACAGACCCGCTG	618
Qy	75	GluLeuTyrLysGlnGlnLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet	94
Db	619	GAGCTGTATACAACAGCGGCGCTCGGGGCGAGCTCCACAGCTCAAGGGCCCTTGACCATG	678
Qy	95	MetAlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGln	114
Db	679	ATGGCCAGCGCACTACACAGGACACTGCGCTCCACCCCGGAAACTTCTCTGTGCAACCCAG	738
Qy	115	IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe	134
Db	739	ATTATCACCTTTGAAAGTTTCAAGAGAACCTGAGGACCTTCTGCTTGTGCATCCCTTT	798
Qy	135	AspCysFtrpGluProValGlnGlu	142
Db	799	GACTGCTGGGAGCCAGTCCAGGAG	822

Search completed: March 11, 2005, 20:27:22  
Job time : 167 secs

GenCore version 5.1.6  
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OM protein - nucleic search, using frame\_plus\_p2n model

Run on: March 11, 2005, 19:37:32 ; Search time 505 Seconds  
(without alignments)  
1672.655 Million cell updates/sec

Title: US-10-723-083-2  
Perfect score: 765  
Sequence: 1 MHHHHHSSGIEGRMAPARS.....ENKDLFLVLPDCEWVQE 142

Scoring table: BLOSUM62

Xgapop 10.0 , Xgapext 0.5  
Ygapop 10.0 , Ygapext 0.5  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 5537552 seqs, 2974263231 residues

Total number of hits satisfying chosen parameters: 11075104

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

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-Q=/cgn2\_1/USPTO.spool/US10723083/runat\_08032005\_131719\_10538/app.query.fasta\_1.327  
-DB=PublishedApplications NA -QFMT=fastap -SUFFIX=rnbp -MINWATCH=0.1  
-LOOPCL=0 -LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=BIOSUM62  
-TRANS=human40.cdi -LIST=45 -DOCALIGN=200 -THR SCORE=pct -THR MAX=100  
-THR\_MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0  
-MAXLEN=200000000 -USER=US10723083@cgn2\_1 1.680 @runat 08032005\_131719\_10538  
-NCPU=6 -ICPU=3 -NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100  
-LONGLOG -DEV TIMEOUT=120 -WARN TIMEOUT=10 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5  
-FGAPOP=6 -FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DLEXT=7

Database :

- Published Applications NA:
- 1: /cgn2\_6/ptodata/1/pubpna/US07\_PUBCOMB.seq.\*
  - 2: /cgn2\_6/ptodata/1/pubpna/PCT\_NEW\_PUB.seq.\*
  - 3: /cgn2\_6/ptodata/1/pubpna/US06\_NEW\_PUB.seq.\*
  - 4: /cgn2\_6/ptodata/1/pubpna/US06\_PUBCOMB.seq.\*
  - 5: /cgn2\_6/ptodata/1/pubpna/US07\_NEW\_PUB.seq.\*
  - 6: /cgn2\_6/ptodata/1/pubpna/PCTUS\_PUBCOMB.seq.\*
  - 7: /cgn2\_6/ptodata/1/pubpna/US08\_NEW\_PUB.seq.\*
  - 8: /cgn2\_6/ptodata/1/pubpna/US08\_PUBCOMB.seq.\*
  - 9: /cgn2\_6/ptodata/1/pubpna/US09A\_PUBCOMB.seq.\*
  - 10: /cgn2\_6/ptodata/1/pubpna/US09B\_PUBCOMB.seq.\*
  - 11: /cgn2\_6/ptodata/1/pubpna/US09C\_PUBCOMB.seq.\*
  - 12: /cgn2\_6/ptodata/1/pubpna/US09\_NEW\_PUB.seq.\*
  - 13: /cgn2\_6/ptodata/1/pubpna/US10A\_PUBCOMB.seq.\*
  - 14: /cgn2\_6/ptodata/1/pubpna/US10B\_PUBCOMB.seq.\*
  - 15: /cgn2\_6/ptodata/1/pubpna/US10C\_PUBCOMB.seq.\*
  - 16: /cgn2\_6/ptodata/1/pubpna/US10D\_PUBCOMB.seq.\*
  - 17: /cgn2\_6/ptodata/1/pubpna/US10E\_PUBCOMB.seq.\*
  - 18: /cgn2\_6/ptodata/1/pubpna/US10F\_PUBCOMB.seq.\*
  - 19: /cgn2\_6/ptodata/1/pubpna/US10\_NEW\_PUB.seq.\*
  - 20: /cgn2\_6/ptodata/1/pubpna/US11\_NEW\_PUB.seq.\*
  - 21: /cgn2\_6/ptodata/1/pubpna/US60\_NEW\_PUB.seq.\*
  - 22: /cgn2\_6/ptodata/1/pubpna/US60\_PUBCOMB.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	765	100.0	429	19	US-10-723-083-3	Sequence 3, Appli
2	765	100.0	458	19	US-10-723-083-1	Sequence 1, Appli
3	682	89.2	777	16	US-10-083-446-55	Sequence 55, Appl
4	678	88.6	402	16	US-10-083-446-176	Sequence 176, App
5	678	88.6	822	16	US-10-083-446-69	Sequence 69, Appl
6	678	88.6	903	16	US-10-083-446-66	Sequence 66, Appl
7	675	88.2	2211	17	US-10-609-346-9	Sequence 9, Appli
8	673	88.0	429	17	US-10-449-831A-141	Sequence 141, App
9	673	88.0	435	9	US-09-826-025-8	Sequence 8, Appli
10	673	88.0	435	14	US-10-083-530-14	Sequence 14, Appl
11	673	88.0	435	17	US-10-188-056-31	Sequence 31, Appl
12	673	88.0	435	17	US-10-188-056-32	Sequence 32, Appl
13	673	88.0	435	17	US-10-411-037-17	Sequence 17, Appl
14	673	88.0	435	17	US-10-411-026-17	Sequence 17, Appl
15	673	88.0	435	17	US-10-410-962-17	Sequence 17, Appl
16	673	88.0	435	17	US-10-411-049-17	Sequence 17, Appl
17	673	88.0	435	18	US-10-410-930-17	Sequence 17, Appl
18	673	88.0	435	18	US-10-410-997-17	Sequence 17, Appl
19	673	88.0	435	18	US-10-411-012-17	Sequence 17, Appl
20	673	88.0	435	18	US-10-287-994-17	Sequence 17, Appl
21	673	88.0	435	18	US-10-410-913-17	Sequence 17, Appl
22	673	88.0	435	18	US-10-785-577-8	Sequence 8, Appli
23	673	88.0	435	19	US-10-410-980-17	Sequence 17, Appl
24	673	88.0	448	17	US-10-609-346-19	Sequence 19, Appl
25	673	88.0	505	18	US-10-688-845-82	Sequence 82, Appl
26	673	88.0	579	17	US-10-449-831A-187	Sequence 187, App
27	673	88.0	781	17	US-10-447-315-20	Sequence 20, Appl
28	673	88.0	789	16	US-10-131-985-16	Sequence 16, Appl
29	673	88.0	789	19	US-10-901-417-16	Sequence 16, Appl
30	673	88.0	1011	13	US-10-044-090-509	Sequence 509, App
31	673	88.0	1318	14	US-10-228-811-3	Sequence 3, Appli
32	673	88.0	1833	9	US-09-783-708-2	Sequence 2, Appli
33	673	88.0	2070	9	US-09-821-883-7	Sequence 7, Appli
34	669	87.5	767	18	US-10-666-122-4	Sequence 4, Appli
35	669	87.5	767	18	US-10-666-122-6	Sequence 6, Appli
36	669	87.5	767	19	US-10-278-698-30	Sequence 30, Appl
37	669	87.5	767	19	US-10-278-698-544	Sequence 544, App
38	668	87.3	381	9	US-09-821-883-19	Sequence 19, Appl
39	668	87.3	435	17	US-10-188-056-33	Sequence 33, Appl
40	668	87.3	435	17	US-10-188-056-34	Sequence 34, Appl
41	668	87.3	496	16	US-10-267-384-191	Sequence 191, App
42	668	87.3	737	15	US-10-081-969-19	Sequence 19, Appl
43	668	87.3	756	15	US-10-177-390-21	Sequence 21, Appl
44	668	87.3	756	17	US-10-351-157-180	Sequence 180, App
45	668	87.3	756	17	US-10-352-554-165	Sequence 165, App

ALIGNMENTS

RESULT 1  
US-10-723-083-3  
; Sequence 3, Application US/10723083  
; Publication No. US20050050602A1  
; GENERAL INFORMATION:  
; APPLICANT: Altosaar, Ilimar  
; APPLICANT: Sardana, Ravinder  
; APPLICANT: Dudani, Aail  
; APPLICANT: Ganz, Peter  
; APPLICANT: Tackaberry, Eileen  
; TITLE OF INVENTION: Production of GM-CSF in Plants  
; FILE REFERENCE: 08-898901US  
; CURRENT APPLICATION NUMBER: US/10/723,083  
; CURRENT FILING DATE: 2003-11-26  
; PRIOR APPLICATION NUMBER: Canada 2,410,702  
; PRIOR FILING DATE: 2002-11-26  
; NUMBER OF SEQ ID NOS: 4  
; SOFTWARE: Patentin version 3.1  
; SEQ ID NO 3  
; LENGTH: 429

TYPE: DNA  
ORGANISM: Homo sapiens  
FEATURE:  
NAME/KEY: CDS  
LOCATION: (1)..(429)  
OTHER INFORMATION:  
US-10-723-083-3

Alignment Scores:  
Pred. No.: 2,88e-92 Length: 429  
Score: 765.00 Matches: 142  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: 19 Gaps: 0

US-10-723-083-2 (1-142) x US-10-723-083-3 (1-429)

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QY 1 MetHisHisHisHisSerSerGlyIleGluGlyArgMetAlaProAlaArgSer 20
DB 1 ATGCACACACACACACACCTCTCCGGCATCGAGGGCGCGATGGCACCGCGCGGTCA 60
QY 21 ProSerProSerThrGlnProTrpGluHisValAsnAlaIleGlnGluAlaArgLeu 40
DB 61 CCAGAGCCCGAGCAGCAGCCCTGGGAGCATGTGAATGCCATCCAGAGGCGCGCGGTCTC 120
QY 41 LeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrValGluValIleSerGlu 60
DB 121 CTGAACTGAGTAGAGACACTGCTGCTGAGATGAATGAACAGTAGAAGTGATATCAGAA 180
QY 61 MetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGluLeuTyrlsGlnGly 80
DB 181 ATGTTTGACCTCCAGAGCGCGACTTGCCTACAGACCGCGCTGGAGCTGTACAGAGCGGC 240
QY 81 LeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMetAlaSerHisTyrls 100
DB 241 CTGCGGGGAGCGCTCAACAGCTCAAGGGCCCTTGACCATGATGTCGCCAGCCACTACAAG 300
QY 101 GlnHisCysProProThrProGluThrSerCysAlaThrGlnIleIleThrPheGluSer 120
DB 301 CAGCATGCGCTCCACCCCGGAACTTCTGTGCAACCGAGATTATCATCTTTGAAAGT 360
QY 121 PheLysGluAsnLeuLysAspPheLeuLeuValIleProPheAspCysTrpGluProVal 140
DB 361 TTCAAAGAGAACCTGAAGGACTTCTGCTGTGTATCCCTTTTGACTGCTGGAGCGCAGTC 420
QY 141 GlnGlu 142
DB 421 CAGGAG 426
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## RESULT 2

US-10-723-083-1  
Sequence 1, Application US/10723083  
Publication No. US20050050602A1  
GENERAL INFORMATION:  
APPLICANT: Altosair, Illimar  
APPLICANT: Sardan, Ravinder  
APPLICANT: Dudani, Aail  
APPLICANT: Ganz, Peter  
APPLICANT: Tackaberry, Eileen  
TITLE OF INVENTION: Production of GM-CSF in Plants  
FILE REFERENCE: 08-89801US  
CURRENT APPLICATION NUMBER: US/10/723,083  
CURRENT FILING DATE: 2003-11-26  
PRIOR APPLICATION NUMBER: Canada 2,410,702  
PRIOR FILING DATE: 2002-11-26  
NUMBER OF SEQ ID NOS: 4  
SOFTWARE: PatentIn version 3.1  
SEQ ID NO 1

LENGTH: 458  
TYPE: DNA  
ORGANISM: Homo sapiens  
FEATURE:

NAME/KEY: CDS  
LOCATION: (10)..(438)  
OTHER INFORMATION:  
US-10-723-083-1

Alignment Scores:  
Pred. No.: 3,15e-92 Length: 458  
Score: 765.00 Matches: 142  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: 19 Gaps: 0

US-10-723-083-2 (1-142) x US-10-723-083-1 (1-458)

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QY 1 MetHisHisHisHisSerSerGlyIleGluGlyArgMetAlaProAlaArgSer 20
DB 10 ATGCACACACACACACACCTCTCCGGCATCGAGGGCGCGATGGCGCGCGCAGC 69
QY 21 ProSerProSerThrGlnProTrpGluHisValAsnAlaIleGlnGluAlaArgLeu 40
DB 70 CCGAGCGCGTCCACACCGCGCTGGAGACACTGAACCGCATCCAGAGGCGCGCGAGGCTC 129
QY 41 LeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrValGluValIleSerGlu 60
DB 130 CTCAACCTCTCCCGCGACACCGCGCGAGATGAACGAGACCGTGGAGGTGATCTCCGAG 189
QY 61 MetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGluLeuTyrlsGlnGly 80
DB 190 ATGTTTGATCTCCAGAGCGCGACTGCTCTCCAGACCGCGCTGTACAGAGCGGC 249
QY 81 LeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMetAlaSerHisTyrls 100
DB 250 CTCCGGGGGAGCGCTCACCAAGCTCAAGGGCCCGCTTCCATGATGGCGTCCCATACAG 309
QY 101 GlnHisCysProProThrProGluThrSerCysAlaThrGlnIleIleThrPheGluSer 120
DB 310 CAGCATGCGCCACCGCGCGAGACCTCTCTCGGCCACCCAGATCATCACCTTCGAGAGC 369
QY 121 PheLysGluAsnLeuLysAspPheLeuLeuValIleProPheAspCysTrpGluProVal 140
DB 370 TTCAGAGGAGAACCTCAAGGACTTCTCTCTCGTATCCCGTTTCGACTGCTGGAGCGGGTG 429
QY 141 GlnGlu 142
DB 430 CAGGAG 435
```

## RESULT 3

US-10-083-446-55  
Sequence 55, Application US/10083446  
Publication No. US20030185790A1  
GENERAL INFORMATION:

APPLICANT: Abrams, Mark A.  
Bauer, S. C.  
Braford-Goldberg, Sarah R.  
Caparon, Mair H.  
Easton, Alan M.  
Klein, Barbara K.  
McKearn, John P.  
Olins, Peter O.  
Paik, Kuman  
Thomas, John W.

TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells  
Using Multivariant (IL-3) Hematopoiesis Chimera Proteins  
NUMBER OF SEQUENCES: 197  
CORRESPONDENCE ADDRESS:

ADDRESSEE: S. Christopher Bauer, Pharmacia Corporation  
Corporate Patent Dept., Mail Zone 04E  
STREET: 800 N. Lindbergh  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63167

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICANT: Abrams, Mark A.  
FILING DATE: 26-Feb-2002  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US/10/083,446  
FILING DATE: 26-Feb-2002  
CLASSIFICATION: <Unknown>  
APPLICATION NUMBER: 08/762,227  
FILING DATE: 09-DEC-1996  
APPLICATION NUMBER: US 08/192,325  
FILING DATE: 14-FEB-1994  
APPLICATION NUMBER: US 08/446,872  
FILING DATE: 06-JUN-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: S. Christopher Bauer  
REGISTRATION NUMBER: 42,305  
REFERENCE/DOCKET NUMBER: C-2790/6  
TELEPHONE: (636)737-6257  
TELEFAX: (636)737-5452  
INFORMATION FOR SEQ ID NO: 55:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 777 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
SEQUENCE DESCRIPTION: SEQ ID NO: 55:  
US-10-083-446-55

Alignment Scores:  
Pred. No.: 8,046-81 Length: 777  
Score: 682.00 Matches: 130  
Percent Similarity: 97.01% Conservative: 0  
Best Local Similarity: 97.01% Mismatches: 4  
Query Match: 89.15% Indels: 0  
DB: 16 Gaps: 0

US-10-723-083-2 (1-142) x US-10-083-446-55 (1-777)

QY 9 SerGlyleGluGlyArgMetAlaProAlaArgSerProSerProSerThrGlnProTrp 28  
DB 376 TCTGGCGGGCTCCAAACATGGCAGCGGCTCTGTTCCCGTCCCGTCTACCCAGCGGTG 435  
QY 29 GluHisValAsnAlaIleGlnGluAlaArgArgLeuAsnLeuSerArgAspThrAla 48  
DB 436 GAACACGTGAATGCCATCCAGGAGGCGCGGCTCTCTGAACCTGAGTAGAGACTGCT 495  
QY 49 AlaGluMetAsnGluThrValGluValIleSerGluMetPheAspLeuGlnProThr 68  
DB 496 GCTGAGATGATGAACAGTAGAGGTATATCAGAAATGTTTACCTCCAGGAGCGACT 555  
QY 69 CysLeuGlnThrArgLeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeu 88  
DB 556 TGCCTACAGACCGCGCTGGAGCTGTACAAGCAGGCGCTGGCGGCGAGCTCACCAAGCTC 615  
QY 89 LysGlyProLeuThrMetMetAlaSerHisTyrLysGlnHisCysProProThrProGlu 108  
DB 616 AAGGGCCCTTTCATGATGAGGACGACCTACCAAGCAGCAGCTGCCCTCCAAACCCCGGAA 675  
QY 109 ThrSerCysAlaThrGlnIleIleThrPheGluSerPheLysGluAsnLeuLysAspPhe 128  
DB 676 ACTTCTGTGCAACCCAGATATATCCTTTGAAGTTTCAAGAGAACCTGAAGGACTTC 735  
QY 129 LeuLeuValIleProPheAspCysTrpGluProValGlnGlu 142  
DB 736 CTGCTTGTATCCCTTTGACTGCTGGAGCCAGTCCAGGAG 777

RESULT 4

US-10-083-446-176

Sequence 176, Application US/10083446  
Publication No. US20030185790A1  
GENERAL INFORMATION:  
APPLICANT: Abrams, Mark A.  
Bauer, S. C.  
Braford-Goldberg, Sarah R.  
Caparon, Mairé H.  
Easton, Alan M.  
Klein, Barbara K.  
McKearn, John P.  
Olin, Peter O.  
Paik, Kumman  
Thomas, John W.  
TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells  
NUMBER OF SEQUENCES: 197  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: S. Christopher Bauer, Pharmacia Corporation  
STREET: 800 N. Lindbergh  
CITY: St. Louis  
STATE: Missouri  
COUNTRY: USA  
ZIP: 63167  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/10/083,446  
FILING DATE: 26-Feb-2002  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/762,227  
FILING DATE: 09-DEC-1996  
APPLICATION NUMBER: US 08/192,325  
FILING DATE: 14-FEB-1994  
APPLICATION NUMBER: US 08/446,872  
FILING DATE: 06-JUN-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: S. Christopher Bauer  
REGISTRATION NUMBER: 42,305  
REFERENCE/DOCKET NUMBER: C-2790/6  
TELEPHONE: (636)737-6257  
TELEFAX: (636)737-5452  
INFORMATION FOR SEQ ID NO: 176:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 402 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
SEQUENCE DESCRIPTION: SEQ ID NO: 176:  
US-10-083-446-176

Alignment Scores:  
Pred. No.: 1,1e-80 Length: 402  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 16 Gaps: 0

US-10-723-083-2 (1-142) x US-10-083-446-176 (1-402)

QY 15 MetAlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
DB 1 ATGGCACCGGCTCGTTCCTCCCGTCTTACCCAGCGGTGGGACACACGTTGATGCCATC 60  
QY 35 GlnGluAlaArgArgLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54

Db 61 CAGGAGGCCGGCGCTCTCTGAACCTGAGTAGACACACTGCTGCTGAGATGAATGAACA 120  
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
Db 121 GTAGAGTGATATCAGAAATGTTTGACCTCAGAGCGGACTTGCCCTACAGACCCGCGCTG 180  
QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 181 GAGCTGTACAAGCAGGCGCTGCGGGGAGCCTCACCAGCTCAAGGGCCCTTGACCATG 240  
QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
Db 241 ATGCCAGCAGCACTACAGAGCAGCACTGCCCTCCACCCCGGAAACTTCTCTGCAACCCAG 300  
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
Db 301 ATTATCACCTTTGAAAGTTTCAAGAGAACTGAGGACTTCTGCTGTGTCATCCCTTT 360  
QY 135 AspCysTrpGluProValGlnGlu 142  
Db 361 GACTGCTGGGAGCAGTCCAGGAG 384

## RESULT 5

US-10-083-446-69

; Sequence 69, Application US/10083446

; Publication No. US20030185790A1

; GENERAL INFORMATION:

; APPLICANT: Abrams, Mark A.

; Bauer, S. C.

; Braford-Goldberg, Sarah R.

; Caparon, Mairé H.

; Easton, Alan M.

; Klein, Barbara K.

; McKearn, John P.

; Olin, Peter O.

; Paik, Kuman

; Thomas, John W.

TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells  
Using Multivariant (IL-3) Hematopoiesis Chimera Proteins

NUMBER OF SEQUENCES: 197

CORRESPONDENCE ADDRESS:

ADDRESSEE: S. Christopher Bauer, Pharmacia Corporation

STREET: 800 N. Lindbergh

CITY: St. Louis

STATE: Missouri

COUNTRY: USA

ZIP: 63167

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent In Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/10/083,446

FILING DATE: 26-Feb-2002

CLASSIFICATION: &lt;Unknown&gt;

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/762,227

FILING DATE: 09-DEC-1996

APPLICATION NUMBER: US 08/192,325

FILING DATE: 14-FEB-1994

APPLICATION NUMBER: US 08/446,872

FILING DATE: 06-JUN-1995

ATTORNEY/AGENT INFORMATION:

NAME: S. Christopher Bauer

REGISTRATION NUMBER: 42,305

REFERENCE/DOCKET NUMBER: C-2790/6

TELECOMMUNICATION INFORMATION:

TELEPHONE: (636)737-6257

TELEFAX: (636)737-5452

INFORMATION FOR SEQ ID NO: 69:

SEQUENCE CHARACTERISTICS:

; LENGTH: 822 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; SEQUENCE DESCRIPTION: SEQ ID NO: 69:  
US-10-083-446-69

## Alignment Scores:

Pred. No.: 2,98e-80 Length: 822  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 16 Gaps: 0

US-10-723-083-2 (1-142) x US-10-083-446-69 (1-822)

QY 15 MetAlaProAlaArgSerProSerProThrGlnProThrGluHisValAsnAlaIle 34  
Db 439 ATGGCACCGGCTCGTTCCTCCCGTCCCGTCTACCCAGCCGTGGACACACGTGAATGCCATC 498  
QY 35 GlnGluAlaArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
Db 499 CAGGAGGCCCGCGCTCTCTGAACTCTAGTAGAGACACTGCTGCTGAGATGAATGAACA 558  
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
Db 559 GTAGAGTGATATCAGAAATGTTTACCTCCAGGAGCCGACTTGCTCTACAGACCCGCGCTG 618  
QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMet 94  
Db 619 GAGCTGTACAAGCAGGCGCTGCGGGGAGCCTCACCAGCTCAAGGGCCCTTGACCATG 678  
QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
Db 679 ATGGCAGCAGCACTACAAGCAGCAGCTGCCCTCCCAACCCCGGAAACTTCTGTGCAACCCAG 738  
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
Db 739 ATTATCACCTTTGAAAGTTTCAAGAGAACTTCTGCTGTGTCATCCCTTT 798  
QY 135 AspCysTrpGluProValGlnGlu 142  
Db 799 GACTGCTGGGAGCAGTCCAGGAG 822

## RESULT 6

US-10-083-446-66

; Sequence 66, Application US/10083446

; Publication No. US20030185790A1

; GENERAL INFORMATION:

; APPLICANT: Abrams, Mark A.

; Bauer, S. C.

; Braford-Goldberg, Sarah R.

; Caparon, Mairé H.

; Easton, Alan M.

; Klein, Barbara K.

; McKearn, John P.

; Olin, Peter O.

; Paik, Kuman

; Thomas, John W.

TITLE OF INVENTION: Methods Of Ex-Vivo Expansion Of Hematopoietic Cells  
Using Multivariant (IL-3) Hematopoiesis Chimera Proteins

NUMBER OF SEQUENCES: 197

CORRESPONDENCE ADDRESS:

ADDRESSEE: S. Christopher Bauer, Pharmacia Corporation

STREET: 800 N. Lindbergh

CITY: St. Louis

STATE: Missouri

COUNTRY: USA

ZIP: 63167

COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/10/083,446  
; FILING DATE: 26-Feb-2002  
; CLASSIFICATION: <Unknown>  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/762,227  
; FILING DATE: 09-DEC-1996  
; APPLICATION NUMBER: US 08/192,325  
; FILING DATE: 14-FEB-1994  
; APPLICATION NUMBER: US 08/446,872  
; FILING DATE: 06-JUN-1995  
; ATTORNEY/AGENT INFORMATION:  
; NAME: S. Christopher Bauer  
; REGISTRATION NUMBER: 42,305  
; REFERENCE/DOCKET NUMBER: C-2790/6  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (636)737-6257  
; TELEFAX: (636)737-5452  
; INFORMATION FOR SEQ ID NO: 66:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 903 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: Double  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; SEQUENCE DESCRIPTION: SEQ ID NO: 66:  
US-10-083-446-66

Alignment Scores:  
Pred. No.: 3,39e-80 Length: 903  
Score: 678.00 Matches: 128  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 88.63% Indels: 0  
DB: 16 Gaps: 0

US-10-723-083-2 (1-142) x US-10-083-446-66 (1-903)

QY 15 MetAlaProAlaArgSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
DB 520 ATGGACCCGGCTGTTCCCGTCCCGTCTACCCAGCCGCGGGAACACGTAATGCCATC 579  
QY 35 GlnGluAlaArgArgLeuLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
DB 580 CAGGAGCCCGCGCTCTCTGAACTCTAGTAGAGACACTGCTGCTGAGATGAATGAACA 639  
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
DB 640 GTAGAAGTGATATCAGAAATGTTTGACCTCCAGAGCCGACTTGCTACAGACCCGCTG 699  
QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrIlysLeuLysGlyProLeuThrMet 94  
DB 700 GAGCTGTACAAGCAGGCGCTGCGGGCAGCGCTCACCAGCTCAAGGGCCCTTGACCATG 759  
QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
DB 760 ATGGCAGCCACATACAAGAGAGCACTGCCCTTCAACCCCGGAACTTCTCTGTGCAACCCAG 819  
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
DB 820 ATTATCACCTTTGAAAGTTTCAAAGAGAACCTGACGACTTCTGCTTGTGTCATCCCTTT 879  
QY 135 AspCysTrpGluProValGlnGlu 142  
DB 880 GACTGTGGGAGCAGTCCAGGAG 903

## RESULT 7

US-10-609-346-9

; Sequence 9, Application US/10609346

; Publication No. US20040063635A1  
; GENERAL INFORMATION:  
; APPLICANT: Yu, Zailin  
; TITLE OF INVENTION: RECOMBINANT HUMAN ALBUMIN FUSION PROTEINS WITH LONG-LASTING BIOLOGICAL ACTIVITY  
; TITLE OF INVENTION: EFFECTS  
; FILE REFERENCE: ZYU-0603  
; CURRENT APPLICATION NUMBER: US/10/609,346  
; PRIOR FILING DATE: 2003-06-26  
; PRIOR APPLICATION NUMBER: US 60/392,948  
; PRIOR FILING DATE: 2002-07-01  
; NUMBER OF SEQ ID NOS: 40  
; SOFTWARE: PatentIn version 3.1  
; SEQ ID NO 9  
; LENGTH: 2211  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: DNA of HSA-GMCSF  
US-10-609-346-9

Alignment Scores:  
Pred. No.: 2,98e-79 Length: 2211  
Score: 675.00 Matches: 127  
Percent Similarity: 100.00% Conservative: 1  
Best Local Similarity: 99.22% Mismatches: 0  
Query Match: 88.24% Indels: 0  
DB: 17 Gaps: 0

US-10-723-083-2 (1-142) x US-10-609-346-9 (1-2211)

QY 15 MetAlaProAlaArgSerProSerThrGlnProTrpGluHisValAsnAlaIle 34  
DB 1825 TTAGCACCCGCGCTGCGCCAGCCGCGGAGCATGTGAATGCCATC 1884  
QY 35 GlnGluAlaArgArgLeuLeuSerArgAspThrAlaAlaGluMetAsnGluThr 54  
DB 1885 CAGGAGCCCGCGCTCTCTGAACTCTAGTAGAGACACTGCTGCTGAGATGAATGAACA 1944  
QY 55 ValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeu 74  
DB 1945 GTAGAAGTGATATCAGAAATGTTTGACCTCCAGAGCCGCGCTACAGACCCGCTG 2004  
QY 75 GluLeuTyrlsGlnGlyLeuArgGlySerLeuThrIlysLeuLysGlyProLeuThrMet 94  
DB 2005 GAGCTGTACAAGCAGGCGCTGCGGGCAGCGCTCACCAGCTCAAGGGCCCTTGACCATG 2064  
QY 95 MetAlaSerHisTyrlsGlnHisCysProProThrProGluThrSerCysAlaThrGln 114  
DB 2065 ATGGCAGCCACATACAAGCAGCACTGCCCTTCAACCCCGGAACTTCTCTGTGCAACCCAG 2124  
QY 115 IleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPhe 134  
DB 2125 ATTATCACCTTTGAAAGTTTCAAAGAGAACCTGACGACTTCTGCTTGTGTCATCCCTTT 2184  
QY 135 AspCysTrpGluProValGlnGlu 142  
DB 2185 GACTGTGGGAGCAGTCCAGGAG 2208

## RESULT 8

US-10-449-831A-141

; Sequence 141, Application US/10449831A

; Publication No. US20040029179A1

; GENERAL INFORMATION:

; APPLICANT: Koenigen, Frank

; TITLE OF INVENTION: Higher molecular weight entities and uses therefor

; FILE REFERENCE: 2385978

; CURRENT APPLICATION NUMBER: US/10/449,831A

; CURRENT FILING DATE: 2003-05-30

; PRIOR APPLICATION NUMBER: USSN 60/384878

; PRIOR FILING DATE: 2002-05-31

; NUMBER OF SEQ ID NOS: 237

; SOFTWARE: PatentIn version 3.2

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; SEQ ID NO 141
; LENGTH: 429
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(429)
; US-10-449-831A-141

Alignment Scores:
Pred. No.: 5,58e-80 Length: 429
Score: 673.00 Matches: 127
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 87.97% Indels: 0
DB: 17 Gaps: 0

US-10-723-083-2 (1-142) x US-10-449-831A-141 (1-429)

QY 16 AlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIleGln 35
Db 49 GCACCGCGCGCTCGCCAGCCGCCAGCAGCAGCCCTGGGAGCATGTGAATGCCATCCAG 108
QY 36 GluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55
Db 109 GAGGCGCGCGCTCTCGAACCTGAGTAGACACTGCTGCTGAGATGAATGAAACAGTA 168
QY 56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75
Db 169 GAAGTCATCTCAGAAATGTTGACCTCCAGGAGCCGACCTGCTCAGACCCCGCTGGAG 228
QY 76 LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95
Db 229 CTGTACAAGCAGGCGCTCGGGGCGGCTCACCAGCTCAAGGGCCCTTGACCATGATG 288
QY 96 AlaSerHisTyrLysGlnHisCysProThrProThrGluThrSerCysAlaThrGlnIle 115
Db 289 GCAGGCACCTACAGCAGCAGCTGCCCTCCAAACCCGGAACCTTCCTGTGCAACCCAGAT 348
QY 116 IleThrPheGluSerPheLysGluAsnLeuLeuLysAspPheLeuValIleProPheAsp 135
Db 349 ATCACCTTTGAAAGTTTCAAGAGAACCTGAGGACTTTCTGCTGTGTCATCCCTTTGAC 408
QY 136 CysTrpGluProValGlnGlu 142
Db 409 TGCTGGAGCCAGTCCAGGAG 429

RESULT 9
US-09-826-025-8
; Sequence 8, Application US/09826025
; Patent No. US20020162123A1
; GENERAL INFORMATION:
; APPLICANT: Chang, Lung-Ji
; TITLE OF INVENTION: Combination Immunogene Therapy
; NUMBER OF SEQUENCES: 25
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Medlen & Carroll, LLP
; STREET: 220 Montgomery Street, Suite 2200
; CITY: San Francisco
; STATE: California
; COUNTRY: United States of America
; ZIP: 94104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA: US/09/826,025
; FILING DATE: 04-Apr-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/838,702
```

```
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Ingolia, Diane E.
; REGISTRATION NUMBER: 40,027
; REFERENCE/DOCKET NUMBER: CHANG-02687
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 705-8410
; TELEFAX: (415) 397-8338
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 435 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "DNA"
; SEQUENCE DESCRIPTION: SEQ ID NO: 8:
US-09-826-025-8

Alignment Scores:
Pred. No.: 5,69e-80 Length: 435
Score: 673.00 Matches: 127
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 87.97% Indels: 0
DB: 9 Gaps: 0

US-10-723-083-2 (1-142) x US-09-826-025-8 (1-435)

QY 16 AlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIleGln 35
Db 52 GCACCGCGCGCTCGCCAGCCGCCAGCAGCAGCCCTGGGAGCATGTGAATGCCATCCAG 111
QY 36 GluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55
Db 112 GAGGCGCGCGCTCTCGAACCTGAGTAGACACTGCTGCTGAGATGAATGAAACAGTA 171
QY 56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75
Db 172 GAAGTCATCTCAGAAATGTTGACCTCCAGGAGCCGACCTGCTCAGACCCCGCTGGAG 231
QY 76 LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95
Db 232 CTGTACAAGCAGGCGCTCGGGGCGGCTCACCAGCTCAAGGGCCCTTGACCATGATG 291
QY 96 AlaSerHisTyrLysGlnHisCysProThrProThrGluThrSerCysAlaThrGlnIle 115
Db 292 GCAGGCACCTACAGCAGCAGCTGCCCTCCAAACCCGGAACCTTCCTGTGCAACCCAGAT 351
QY 116 IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPheAsp 135
Db 352 ATCACCTTTGAAAGTTTCAAGAGAACCTGAGGACTTTCTGCTGTGTCATCCCTTTGAC 411
QY 136 CysTrpGluProValGlnGlu 142
Db 412 TGCTGGAGCCAGTCCAGGAG 432

RESULT 10
US-10-083-590-14
; Sequence 14, Application US/10083590
; Publication No. US20030027257A1
; GENERAL INFORMATION:
; APPLICANT: IATROU, Kostas
; APPLICANT: FARRELL, Patrick J.
; APPLICANT: BEHIE, Leo A.
; TITLE OF INVENTION: SEQUENCES FOR IMPROVING THE EFFICIENCY OF SECRETION OF
; FILE REFERENCE: 028722-207
; CURRENT APPLICATION NUMBER: US/10/083,590
; CURRENT FILING DATE: 2002-02-27
; PRIOR APPLICATION NUMBER: EARLIER FILING DATE: 1999-02-24
; PRIOR FILING DATE: EARLIER FILING DATE: 1999-02-24
; APPLICATION NUMBER: EARLIER APPLICATION NUMBER: US 09/136,421
```



; PRIOR FILING DATE: EARLIER FILING DATE: 1998-08-20  
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: US 60/056,871  
; PRIOR FILING DATE: EARLIER FILING DATE: 1997-08-21  
; NUMBER OF SEQ ID NOS: 14  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 14  
; LENGTH: 435  
; TYPE: DNA  
; ORGANISM: Human  
US-10-083-590-14

Alignment Scores:  
Pred. No.: 5,69e-80 Length: 435  
Score: 673.00 Matches: 127  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 87.97% Indels: 0  
DB: 14 Gaps: 0

US-10-723-083-2 (1-142) x US-10-083-590-14 (1-435)

QY 16 AlaProAlaArgSerProSerProSerThrGlnProTTPGluHisValAlaAlaGln 35  
DB 52 GCACCGCGCGCTCGCCAGCCCGCAGCAGCGCCCTGGGAGCATGTGAATGCCATCCAG 111  
QY 36 GluAlaArgLeuLeuLeuLeuLeuSerA:GAspThrAlaAlaGluMetAenGluThrVal 55  
DB 112 GAGGCCGCGCGTCTCTGAACTGTAGTAGACACTGCTGCTGAGATGAATGAACAGTA 171  
QY 56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75  
DB 172 GAAGTCATCTCAGAAATGTTTACCTCCAGGAGCGACCTGACAGACCCGCGCTGGAG 231  
QY 76 LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95  
DB 232 CTGTACAGCAGCGGCTCGGGGCGAGCTCACCAGCTCAAGGCCCTTGACCATGATG 291  
QY 96 AlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIle 115  
DB 292 GCACGCGCACTACAAGCAGCACTGCCCTCCAAACCCCGGAACTTCTGTGTGAACCCAGATT 351  
QY 116 IleThrPheGluSerPheLysGluAenLeuLeuLeuLeuLeuLeuLeuValIleProPheAsp 135  
DB 352 ATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTTCTGCTGTGATCCCTTTGAC 411  
QY 136 CysTTPGluProValGlnGlu 142  
DB 412 TGCTGGAGCCAGTCCAGGAG 432

RESULT 11

US-10-188-056-31  
; Sequence 31, Application US/10188056  
; Publication No. US20040009934A1  
; GENERAL INFORMATION:  
; APPLICANT: Qiu, Jian-Tai  
; APPLICANT: Lai, Wan-Ching  
; APPLICANT: Chu, Yong Liang  
; APPLICANT: Li, Frank Q.  
; TITLE OF INVENTION: Improved GM-CSF Nucleic Acid Sequences  
; FILE REFERENCE: 3781-004-27  
; CURRENT APPLICATION NUMBER: US/10/188,056  
; CURRENT FILING DATE: 2002-09-26  
; NUMBER OF SEQ ID NOS: 34  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 31  
; LENGTH: 435  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-188-056-31

Alignment Scores:  
Pred. No.: 5,69e-80 Length: 435  
Score: 673.00 Matches: 127

Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 87.97% Indels: 0  
DB: 17 Gaps: 0

US-10-723-083-2 (1-142) x US-10-188-056-31 (1-435)

QY 16 AlaProAlaArgSerProSerProSerThrGlnProTTPGluHisValAlaAlaGln 35  
DB 52 GCTCCCGCCAGAACCCAGCCCTCCACCCAGCCCTGGGAGCACGTGAACGCCATCCAG 111  
QY 36 GluAlaArgLeuLeuLeuLeuLeuSerA:GAspThrAlaAlaGluMetAenGluThrVal 55  
DB 112 GAGGCCAGACGGCTGCTGAACCTGTCCAGAGACCCGCCGCGAGATGAACGAGACCGTG 171  
QY 56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75  
DB 172 GAGGTGATCAGCGAGATGTTTCGACCTGCAGAGGCCACCTGCTGTCAGACCCGCGCTGGAG 231  
QY 76 LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95  
DB 232 CTGTACAGCAGCGGCTCGGGGCGAGCTGACCAAGCTGAAGGGACCCCTGACCATGATG 291  
QY 96 AlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIle 115  
DB 292 GCACGCGCACTACAAGCAGCACTGCCCTCCACACCCCGAGACGCTGCCGCCACCCAGATC 351  
QY 116 IleThrPheGluSerPheLysGluAenLeuLeuLeuLeuLeuLeuValIleProPheAsp 135  
DB 352 ATCACCTTCGAGAGCTTCAAGGAGAACCTGAAGGACTTCTGCTGTGATCCCTTTGAC 411  
QY 136 CysTTPGluProValGlnGlu 142  
DB 412 TGCTGGAGCCCGTCCAGGAG 432

RESULT 12

US-10-188-056-32  
; Sequence 32, Application US/10188056  
; Publication No. US20040009934A1  
; GENERAL INFORMATION:  
; APPLICANT: Qiu, Jian-Tai  
; APPLICANT: Lai, Wan-Ching  
; APPLICANT: Chu, Yong Liang  
; APPLICANT: Li, Frank Q.  
; TITLE OF INVENTION: Improved GM-CSF Nucleic Acid Sequences  
; FILE REFERENCE: 3781-004-27  
; CURRENT APPLICATION NUMBER: US/10/188,056  
; CURRENT FILING DATE: 2002-09-26  
; NUMBER OF SEQ ID NOS: 34  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 32  
; LENGTH: 435  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-188-056-32

Alignment Scores:  
Pred. No.: 5,69e-80 Length: 435  
Score: 673.00 Matches: 127  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 87.97% Indels: 0  
DB: 17 Gaps: 0

US-10-723-083-2 (1-142) x US-10-188-056-32 (1-435)

QY 16 AlaProAlaArgSerProSerProSerThrGlnProTTPGluHisValAlaAlaGln 35  
DB 52 GCACCGCGCGCTCGCCAGCCCGCAGCAGCCCTGGGAGCATGTGAATGCCATCCAG 111  
QY 36 GluAlaArgLeuLeuLeuLeuLeuSerA:GAspThrAlaAlaGluMetAenGluThrVal 55  
DB 112 GAGGCCGCGCGTCTCTGAACTGTAGTAGACACTGCTGCTGATGAATGAACAGTA 171

Qy	56	GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu	75
Db	172	GAAAGTCATCTCAGAAATGTTTGACCTCCAGGACCGAGCTGCTACAGACCGCTCGAG	231
Qy	76	LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet	95
Db	232	CTGTACACAGAGGGCTTGGGGGACGCTCACAGCTCAAGGGCCCTTGACCATGATG	291
Qy	96	AlaSerHisTyrLysGlnHisCysProThrProGluThrSerCysAlaThrGlnIle	115
Db	292	GCCAGACCACTACAAAACAGCACTGCCCTCCAAACCCGGAAACTTCCTGTGCACACAGATT	351
Qy	116	IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPheAsp	135
Db	352	ATCACCTTGTAAAGTTTCAAGAGAACCTGAAGGACTTTCGTGTGTCATCCCCCTTTGAC	411
Qy	136	CysTrpGluProValGlnGlu	142
Db	412	TGCTGGGAGCCAGTCACAGAG	432

Qy	36	GluAlaArgLeuLeuAsnLeuSerArgAspThrIaLaGluMetAsnGluThrVal	55
Db	112	GAGGCCCGCGCTCCTGAACCTGAGTAGACACTGCTGTGAGATGAATGAACAGTA	171
Qy	56	GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu	75
Db	172	GAAGTCATCTCAGAATAATTGTTGACCTCCAGAGCGGACCCTGCCTACAGACCCGCTTGGAG	231
Qy	76	LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysIleuLysGlyProLeuThrMetMet	95
Db	232	CTGTACAACAGGGCTCGGGGCGAGCTCACCAAGCTCAAGGGCCCCCTTGACCATGATG	291
Qy	96	AlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIle	115
Db	292	GCCAGCCACTACAAGCAGCACTGCGCTCCAACCCCGGAAATCTCTGTGCAACCCAGATT	351
Qy	116	IleThrPheGluSerPheLysGluAsnLeuLysAspPheIleuValIleProPheAsp	135
Db	352	ATCACCTTTGAAGAAGTTTCAAAGAGAACCTGAGAGACTTTCGTCTGTGCATCCCCTTTGAC	411
Qy	136	CysTrpGluProValGlnGlu	142
Db	412	TGCTGGAGCCAGTCCAGGAG	432

QY 16 AlaProAlaArgSerProSerProSerThrGlnProThrGluHisValAsnAlaIleGln 35  
DB 52 GCACCCGCGCGTCTGCGCCAGCCAGCAGCAGCCCTGGAGCATGTGAATGCCATCCAG 111  
QY 36 GluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55  
DB 112 GAGGCCGCGCGTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAATGAACAGTA 171  
QY 56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75  
DB 172 GAAGTCATCTCAGAAATGTTGACCTCCAGGAGCCGACCTGCCTACAGACCCGCGCTGGAG 231  
QY 76 LeuTyrLysGlnGluLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95  
DB 232 CTGTACAGCAGGCGCTGCGGGCAGCCCTCACAAGCTCAAGGGCCCTTGACCATGATG 291  
QY 96 AlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIle 115  
DB 292 GCCAGCCTACNAGCAGCACTGCCCTCCAAACCCCGGAACTTCTGTGCAACCCAGATT 351  
QY 116 IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPheAsp 135  
DB 352 ATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTTCTGCTGTGTCATCCCTTTGAC 411  
QY 136 CysTyrGluProValGlnGlu 142  
DB 412 TGCTGGGAGCCAGTCCAGGAG 432

## RESULT 15

US-10-410-962-17  
; Sequence 17, Application US/10410962  
; Publication No. US2004007836A1  
; GENERAL INFORMATION:  
; APPLICANT: Neose Technologies, Inc.  
; APPLICANT: DeFrees, Shawn  
; APPLICANT: Zopf, David  
; APPLICANT: Bayer, Robert  
; APPLICANT: Hakes, David  
; APPLICANT: Chen, Xi  
; APPLICANT: Bowe, Caryn  
; TITLE OF INVENTION: GRANULOCYTE COLONY STIMULATING FACTOR: REMODELING AND  
; TITLE OF INVENTION: GLYCOCONJUGATION OF G-CSF  
; FILE REFERENCE: 040853-01-5054  
; CURRENT APPLICATION NUMBER: US/10/410,962  
; CURRENT FILING DATE: 2003-04-09  
; PRIOR APPLICATION NUMBER: US 60/328,523  
; PRIOR FILING DATE: 2001-10-10  
; PRIOR APPLICATION NUMBER: US 60/344,692  
; PRIOR FILING DATE: 2001-10-19  
; PRIOR APPLICATION NUMBER: US 60/387,292  
; PRIOR FILING DATE: 2002-06-07  
; PRIOR APPLICATION NUMBER: US 60/391,777  
; PRIOR FILING DATE: 2002-06-25  
; PRIOR APPLICATION NUMBER: US 60/396,594  
; PRIOR FILING DATE: 2002-07-17  
; PRIOR APPLICATION NUMBER: US 60/404,249  
; PRIOR FILING DATE: 2002-08-16  
; PRIOR APPLICATION NUMBER: US 60/407,527  
; PRIOR FILING DATE: 2002-08-28  
; NUMBER OF SEQ ID NOS: 75  
; SOFTWARE: Patent in version 3.2  
; SEQ ID NO 17  
; LENGTH: 435  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-410-962-17

Alignment Scores:  
Pred. No.: 5,69e-80 Length: 435  
Score: 673.00 Matches: 127  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 87.97% Indels: 0

DB: 17 Gaps: 0  
US-10-723-083-2 (1-142) x US-10-410-962-17 (1-435)  
QY 16 AlaProAlaArgSerProSerProSerThrGlnProThrGluHisValAsnAlaIleGln 35  
DB 52 GCACCCGCGCGTCTGCGCCAGCCAGCAGCAGCCCTGGAGCATGTGAATGCCATCCAG 111  
QY 36 GluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55  
DB 112 GAGGCCGCGCGTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAATGAACAGTA 171  
QY 56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75  
DB 172 GAAGTCATCTCAGAAATGTTGACCTCCAGGAGCCGACCTGCCTACAGACCCGCGCTGGAG 231  
QY 76 LeuTyrLysGlnGluLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95  
DB 232 CTGTACAGCAGGCGCTGCGGGCAGCCCTCACAAGCTCAAGGGCCCTTGACCATGATG 291  
QY 96 AlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIle 115  
DB 292 GCCAGCCTACNAGCAGCACTGCCCTCCAAACCCCGGAACTTCTGTGCAACCCAGATT 351  
QY 116 IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPheAsp 135  
DB 352 ATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTTCTGCTGTGTCATCCCTTTGAC 411  
QY 136 CysTyrGluProValGlnGlu 142  
DB 412 TGCTGGGAGCCAGTCCAGGAG 432

Search completed: March 11, 2005, 21:35:24  
Job time : 512 secs



GenCore version 5.1.6  
Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - nucleic search, using frame\_plus\_p2n model

Run on: March 11, 2005, 18:16:26 ; Search time 2822 Seconds  
(without alignments)  
1915.353 Million cell updates/sec

Title: US-10-723-083-2

Perfect score: 765

Sequence: 1 MHHHHHSSGIERMAPARS.....ENLKDFLLVDFCWPQVE 142

Scoring table:

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-DB=EST -QFMT=fastap -SUFFIX=rst -MINMATCH=0.1 -LOOPEXT=0 -LOOPEXT=0  
-UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45  
-DOCALIGN=200 -THR\_SCORE=pct -THR\_MAX=100 -THR\_MIN=0 -ALIGN=15 -MODE=LOCAL  
-OUTFMT=pcio -NORM=ext -HEAPSIZ=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US10723083 @CGN 1.1 4352 @runat.08032005.131716.10436 -NCPU=6 -ICPU=3  
-NO WMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG  
-DSV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPOP=6  
-FGAPEXT=7 -XGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

-MODEL=frame+ p2n.model -DEV=xlh  
-O=/cgn2.1/USPTO.spool-US10723083/runat.08032005.131716.10436/app.query.fasta\_1.327  
-DB=EST -QFMT=fastap -SUFFIX=rst -MINMATCH=0.1 -LOOPEXT=0 -LOOPEXT=0  
-UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45  
-DOCALIGN=200 -THR\_SCORE=pct -THR\_MAX=100 -THR\_MIN=0 -ALIGN=15 -MODE=LOCAL  
-OUTFMT=pcio -NORM=ext -HEAPSIZ=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US10723083 @CGN 1.1 4352 @runat.08032005.131716.10436 -NCPU=6 -ICPU=3  
-NO WMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG  
-DSV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPOP=6  
-FGAPEXT=7 -XGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

EST:  
1: gb\_est1:\*  
2: gb\_est2:\*  
3: gb\_hic:\*  
4: gb\_est3:\*  
5: gb\_est4:\*  
6: gb\_est5:\*  
7: gb\_est6:\*  
8: gb\_gsal:\*  
9: gb\_g882:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	673	88.0	588	2	AW207707 UI-H-B12-
2	673	88.0	658	5	EX111836 EX111836
3	673	88.0	660	2	BE218982 hv47a07.x
4	673	88.0	666	1	AI912784 we13f07.x
5	673	88.0	672	2	BE671554 7e53h07.x
6	673	88.0	695	2	BE669962 7e27g08.x
7	673	88.0	895	2	BE873976 601484045
8	668	87.3	584	7	CF341802 TGESTzyj4
9	652	85.2	592	7	CF370966 TGESTzyj5

10	647	84.6	565	2	BF938995
11	647	84.6	585	7	CF370833
12	640	83.7	718	6	CD369973
13	614	80.3	701	5	BU633411
14	563	73.6	666	6	CA307828
15	559	73.1	475	1	AA995402
16	534	69.8	661	6	CD368851
17	480.5	62.8	561	6	CB457551
18	480.5	62.8	672	7	CF614774
19	473	61.8	572	4	BM539160
20	469.5	61.4	511	6	CB430266
21	458	59.9	336	2	AM951121
22	433	59.2	608	6	CD367244
23	419	54.8	423	2	AW784714
24	402	52.5	549	1	AI677936
25	375	49.0	531	4	BG236310
26	343	44.8	274	1	AA361936
27	301.5	39.4	647	2	BB533718
28	269.5	35.2	629	2	BB664267
29	268	35.0	269	7	CF370885
30	260	34.0	517	6	CB430980
31	254	33.2	369	5	BX521029
32	248	32.4	483	1	AI180669
33	245	32.0	244	4	BG236058
34	229	29.9	470	1	AI121878
35	214	28.0	697	9	AG112609
36	171	22.4	160	7	CF341980
37	162	21.2	136	7	CF341168
38	157	20.5	892	9	CR059250
39	102	13.3	935	7	CF264246
40	101	13.2	721	8	AZ969504
41	99.5	13.0	131	8	AF179193
42	98	12.8	372	7	T29160
43	94	12.3	828	7	CK144209
44	92	12.0	405	5	BU865195
45	91.5	12.0	699	1	AJ819401

#### ALIGNMENTS

RESULT 1  
AW207707

LOCUS  
DEFINITION

UI-H-B12-age-e-09-0-UI.s1 NCI\_CGAP\_Sub4 Homo sapiens cDNA clone  
IMAGE:2724184 3', mRNA sequence.

ACCESSION  
AW207707

VERSION  
AW207707.1 GI:6507203

KEYWORDS  
EST.

SOURCE  
Homo sapiens (human)

ORGANISM  
Homo sapiens

REFERENCE  
1 (bases 1 to 588)

AUTHORS  
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

TITLE  
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

COMMENT  
Unpublished (1997)

CONTACT  
Contact: Robert Strausberg, Ph.D.

EMAIL  
Email: cgapsb@mail.nih.gov

OLIGO-DT  
Oligo-dT track not found, Not I site shown in beginning of sequence

IS  
is likely internal to the message. cDNA library preparation: M.B.

SCORES  
Scores lab clone distribution: NCI-CGAP clone distribution

INFORMATION  
Information can be found through the I.M.A.G.E. Consortium/LLNL at:

WWW-BIO-ILLN-  
www-bio-illn.gov/bbrp/image/image.html

SEQ PRIMER  
Seq primer: M13 Forward

POLYA-NO  
POLYA-NO. Location/Qualifiers

FEATURES  
source

1..588

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="IMAGE:2724184"

/lab\_host="DH10B (Life Technologies)"



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Db      559 GAGGCCGGCGTCTCTGAACTGAGTACAGACACTGCTGCTGAGATGAATGAACAGTA 500
Qy      56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75
Db      499 GAACTCATCTCAGAAATGTTTGACCTCCAGGAGCCGACCTGCCTACAGACCCGCTGGAG 440
Qy      76 LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95
Db      439 CTGTACAAGCAGGGCTCGCGGCGACCTCACCAGCTCAAGGGCCCTTGACCATGATG 380
Qy      96 AlaSerHisTyrLysGlnHisCysProThrProGluThrSerCysAlaThrGlnIle 115
Db      379 GCCAGCATTACAGAGCAGCACTGCCCTCAACCCCGGAACTTCTCTGTGCAACCCAGATT 320
Qy      116 IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPheAsp 135
Db      319 ATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTTCTGCTGTGTCATCCCTTTGAC 260
Qy      136 CysTrpGluProValGlnGlu 142
Db      259 TCGTGGAGCCAGTCCAGGAG 239

RESULT 3
LOCUS      BE218982
DEFINITION hv47a07.x1 NCI CGAP Lu24 Homo sapiens cDNA clone IMAGE:3176532 3'
            similar to gb:M11220 GRANULOCYTE-MACROPHAGE COLONY-STIMULATING
            FACTOR PRECURSOR (HUMAN);, mRNA sequence.
ACCESSION  BE218982
VERSION     BE218982.1 GI:8906300
KEYWORDS    EST.
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
            1 (bases 1 to 660)
REFERENCE   NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS     National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL     Unpublished (1997)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgabbs-remail.nih.gov
            Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
            Emmert-Buck, M.D., Ph.D.
            cDNA Library Preparation: M. Bento Soares, Ph.D.
            DNA Sequencing by: Greg Lennon, Ph.D.
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LNL, send email to:
            info@image.llnl.gov
            Seq primer: -40UP from Gibco
            High quality sequence stop: 445.
FEATURES    Location/Qualifiers
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            /organism="Homo sapiens"
            /mol_type="mRNA"
            /db_xref="taxon:9606"
            /clone="IMAGE:3176532"
            /tissue_type="carcinoid"
            /lab_host="DH10B"
            /clone_lib="NCI CGAP Lu24"
            /notes="Organ: lung; Vector: pTT73D-Pac (Pharmacia) with a
            modified polylinker; plasmid DNA from the normalized
            library NCI CGAP Lu5 was prepared, and ss circles were
            made in vitro. Following HAP purification, this DNA was
            used as tracer in a subtractive hybridization reaction.
            The driver was PCR-amplified cDNAs from a pool of 5,000
            clones made from the same library (cloneIDs
            1414920-1417991 and 1520904-1522439). Subtraction by Bento
            Soares and M. Fatima Bonaldo."
ORIGIN
Alignment Scores:

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Pred. No.:      6,88e-63      Length:      660
Score:          673.00      Matches:      127
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match:    87.97%      Indels:      0
DB:            2      Gaps:      0

US-10-723-083-2 (1-142) x BE218982 (1-660)

Qy      16 AlaProAlaArgSerProSerThrGlnProTrpGluHisValAsnAlaIleGln 35
Db      56 GCACCCGCGCTCGCCAGACCCAGCAGCGAGCCCTGGGAGCATGTGAATGCCATCCAG 115
Qy      36 GluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55
Db      116 GAGGCCCGCGCTCTCTGAACCTGAGTACAGACACTGCTGCTGAGATGAATGAACAGTA 175
Qy      56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75
Db      176 GAAATCATCTCAGAAATGTTTACCTCCAGGAGCCGACCTGCTCAGACCCGCTGGAG 235
Qy      76 LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95
Db      236 CTGTACAAGCAGGGCTCGCGGCGACCTCACCAGCTCAAGGGCCCTTGACCATGATG 295
Qy      96 AlaSerHisTyrLysGlnHisCysProThrProGluThrSerCysAlaThrGlnIle 115
Db      296 GCCAGCATTACAGAGCAGCACTGCCCTCAACCCCGGAACTTCTCTGTGCAACCCAGATT 355
Qy      116 IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPheAsp 135
Db      356 ATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTTCTGCTGTGTCATCCCTTTGAC 415
Qy      136 CysTrpGluProValGlnGlu 142
Db      416 TCGTGGAGCCAGTCCAGGAG 436

RESULT 4
LOCUS      AI912784
DEFINITION AI912784
            similar to gb:M11220 GRANULOCYTE-MACROPHAGE COLONY-STIMULATING
            FACTOR PRECURSOR (HUMAN);, mRNA sequence.
ACCESSION  AI912784
VERSION     AI912784.1 GI:5632639
KEYWORDS    EST.
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
            1 (bases 1 to 666)
REFERENCE   NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS     National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL     Unpublished (1997)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgabbs-remail.nih.gov
            Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
            Emmert-Buck, M.D., Ph.D.
            cDNA Library Preparation: M. Bento Soares, Ph.D.
            DNA Sequencing by: Greg Lennon, Ph.D.
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LNL at:
            www-bio.llnl.gov/bbrp/image/image.html
            Insert length: 743
            Seq primer: -40UP from Gibco
            High quality sequence stop: 447.
FEATURES    Location/Qualifiers
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            /organism="Homo sapiens"
            /mol_type="mRNA"
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ACCESSION      BE669962
VERSION        BE669962.1  GI:10030503
KEYWORDS      EST.
SOURCE        Homo sapiens (human)
ORGANISM      Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 695)
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE        National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
              Tumor Gene Index
JOURNAL        Unpublished (1997)
COMMENT      Contact: Robert Strausberg, Ph.D.
              Email: cgapbs-remail.nih.gov
              Emmert-Buck, M.D., Ph.D.
              cDNA Library Prepared by: Greg Lennon, Ph.D.
              DNA Sequencing by: Washington University Genome Sequencing Center
              Clone distribution: NCI-CGAP clone distribution information can be
              found through the I.M.A.G.E. Consortium/LINL, send email to:
              info@image.llnl.gov
              Seq primer: -40UP from Gibco
              High quality sequence stop: 456.
FEATURES      Location/Qualifiers
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                      /mol_type="mRNA"
                      /db_xref="taxon:9606"
                      /clone="IMAGE:3283742"
                      /tissue_type="carcinoid"
                      /lab_host="DH10B"
                      /clone_lib="NCI CGAP Lu24"
                      /notes="Organ: lung; Vector: pT7T3D-Pac (Pharmacia) with a
                      modified polylinker; plasmid DNA from the normalized
                      library NCI CGAP Lu5 was prepared, and ss circles were
                      made in vitro. Following HAP purification, this DNA was
                      used as tracer in a subtractive hybridization reaction.
                      The driver was PCR-amplified cDNAs from a pool of 5,000
                      clones made from the same library (cloneIDs
                      1414920-1417991 and 1520904-1522439). Subtraction by Bento
                      Soares and M. Fatima Bonaldo."
ORIGIN
Alignment Scores:
Pred. No.:      7.38e-63      Length:      695
Score:          673.00      Matches:     127
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match:    87.97%      Indels:     0
DB:             2      Gaps:       0

US-10-723-083-2 (1-142) x BE669962 (1-695)
QY      16 AlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIleGln 35
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QY      36 GluAlaArgGluLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55
Db      152 GAGGCCGCCGCTCTCTGAACCTGTAGTACACACTGCTGCTGAGATGAATGAACAGTA 211
QY      56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75
Db      212 GAAGTCATCTCAGAAATGTTTGACCTCCAGGACCGACCTGCTCAGACCCGCCCTGGAG 271
QY      76 LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95
Db      272 CTGTACAAGCAGGCGCTCGGGGGCAGCCTCACCAGCTCAAGGGCCCTTGACCATGATG 331
QY      96 AlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIle 115

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Db      332 GCAGGCCACTACAAGCAGCACTGCCCTCCAAACCCCGAAACTTCCTGTGCAACCCAGATT 391
QY      116 IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPheAsp 135
Db      392 ATCACCTTTGAAAGTTTCAAGAGAACCTGAAGGACTTTCTGCTGTGTCATCCCTTTGAC 451
QY      136 CysTrpGluProValGlnGlu 142
Db      452 TGCTGGGAGCCAGTCAGTCAGGAG 472

RESULT 7
LOCUS      BE873976
DEFINITION BE873976
ACCESSION BE873976
VERSION   BE873976.1  GI:10322752
KEYWORDS  EST.
SOURCE    Homo sapiens (human)
ORGANISM  Homo sapiens
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 895)
AUTHORS  NIH-MGC http://mgc.nci.nih.gov/
TITLE    National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL  Unpublished (1999)
COMMENT  Contact: Robert Strausberg, Ph.D.
          Email: cgapbs-remail.nih.gov
          Tissue Procurement: DCTD/Drp/Gazdar
          cDNA Library Preparation: Life Technologies, Inc.
          cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LINL)
          DNA Sequencing by: Incyte Genomics, Inc.
          Clone distribution: MGC clone distribution information can be
          found through the I.M.A.G.E. Consortium/LINL at:
          http://image.llnl.gov
          Plate: L1AM9663 row: j column: 12
          High quality sequence stop: 711.
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                  /mol_type="mRNA"
                  /db_xref="taxon:9606"
                  /clone="IMAGE:3886571"
                  /tissue_type="large cell carcinoma, undifferentiated"
                  /lab_host="DH10B (phage-resistant)"
                  /clone_lib="NIH MGC 69"
                  /note="Organ: lung; Vector: pCMV-SPORT6; Site 1: NotI;
                  Site 2: SalI; Cloned unidirectionally. Primer: Oligo dT.
                  Average insert size 1.1 kb. Library constructed by Life
                  Technologies."
ORIGIN
Alignment Scores:
Pred. No.:      1.03e-62      Length:      895
Score:          673.00      Matches:     127
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match:    87.97%      Indels:     0
DB:             2      Gaps:       0

US-10-723-083-2 (1-142) x BE873976 (1-895)
QY      16 AlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIleGln 35
Db      50 GCACCCGCCGCTCGCCAGCCCGCCAGCAGCCAGCCCTGGGAGCATGTGAATGCATCCAG 109
QY      36 GluAlaArgGluLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55
Db      110 GAGGCCGCCGCTCTCTGAACCTGTAGTACACACTGCTGCTGAGATGAATGAACAGTA 169
QY      56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75
Db      170 GAAGTCATCTCAGAAATGTTTGACCTCCAGGAGCCGACCTGCTCAGACCCGCCCTGGAG 229

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76 LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95  
 230 CTGTACAAGCAGGCGCTCGGGGAGGCTACCAAGCTCAAGGCGCCCTTGACCATGATG 289  
 96 AlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIle 115  
 290 GCCAGCCACTACAAGCAGCAGCTGCGCTCCAACCGCGAAATTCCTGTGCAACCCAGATT 349  
 116 IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPheAsp 135  
 350 ATCACCTTTGAAGATTTCAAGAGAACTGAAGACTTTCTGCTGTGTCATCCCTTTGAC 409  
 136 CysTrpGluProValGlnGlu 142  
 410 TGCTGGGAGCCAGTCCAGGAG 430

RESULT 8  
 CF341802  
 LOCUS  
 DEFINITION  
 TgESTyJ43f02.y1 Tg CAST Tachyzoite cDNA library Toxoplasma gondii  
 cDNA clone TgESTyJ43f02.y1 5' similar to SW:CSF2 HUMAN P04141  
 GRANULOCYTE-MACROPHAGE COLONY-STIMULATING FACTOR PRECURSOR ; mRNA  
 sequence.  
 CF341802  
 CF341802.1 GI:33831915  
 EST.  
 Toxoplasma gondii  
 Toxoplasma gondii  
 Toxoplasma gondii  
 Sarcocystidae; Toxoplasma.  
 1 (bases 1 to 584)  
 Clifton,S., Page,D., Martin,J., Wyllie,T., Dante,M., Marra,M.,  
 Hillier,L., Kucaba,T., Theising,B., Bowers,Y., Gibbons,M.,  
 Ritter,E., Bennett,J., Franklin,C., Tsagarelisvili,R., Ronko,I.,  
 Kennedy,S., Maguire,L., Waterston,R. and Wilson,R.  
 Toxoplasma EST Project  
 Unpublished (2001)  
 Contact: Clifton, S.  
 Toxoplasma EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: toxo@wustl.edu  
 Contact David Sibley (toxost@borcim.wustl.edu) for further  
 information relating to organism, libraries, or clone availability.  
 Seq primer: -400P from Gibco.

FEATURES  
 source  
 1..584  
 Location/Qualifiers  
 /organism="Toxoplasma gondii"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:5811"  
 /clone="TgESTyJ43f02.y1"  
 /dev\_stage="Tachyzoite"  
 /lab\_host="ElectroTen Blue cells (Stratagene)"  
 /clone\_lib="Tg CAST Tachyzoite cDNA Library"  
 /notes="Vector: Modified pBluescript (pBS SK+); Site\_1:  
 BamHI; Site\_2: EcoRI; The cDNA library was constructed by  
 Keliang Tang, and Robert Cole at Washington University.  
 cDNA was synthesized from poly(A)+ mRNA using the  
 template-switching PCR method (SMART cDNA Kit, BD  
 Biosciences). First strand cDNA was reverse transcribed  
 using the CDS III/3' primer and a 5' template switch  
 primer (Smart IV primer). The product of the first strand  
 synthesis was PCR amplified using the same primer set and  
 the fragments were digested with SfiI. The fragments were  
 size selected, ligated into a modified pBluescript vector  
 (obtained from Michael White, Montana State University)  
 containing directional SfiI sites, and electroporated into  
 ElectroTen Blue cells. Vector: SfiI sites were added to  
 the multiple cloning region of pBluescript SK+ between the

BamHI/EcoRI sites. The modified polylinker has the  
 following sequence: 5'GAATTCGGCCATTACGGCC(G)n- insert--  
 GCGCCCTCGCCACGATCC3' where n=3-4 G nucleotides.  
 WARNING: the library contains a small percentage of cDNAs  
 derived from the human host cells. Library materials  
 provided by David Sibley, Washington University."

## ORIGIN

Alignment Scores: 2.05e-62 Length: 584  
 Pred. No.: 668.00 Matches: 126  
 Score: 99.21% Conservative: 0  
 Percent Similarity: 99.21% Mismatches: 1  
 Best Local Similarity: 99.21% Indels: 0  
 Query Match: 87.32% Gaps: 0  
 DB: 7

US-10-723-083-2 (1-142) x CF341802 (1-584)

QY 16 AlaProAlaArgSerProSerProThrGlnProTrrGluHisValAsnAlaIleGln 35  
 Db 84 GCACCGCCCGCTCGCCAGCCCGCCAGCGAGCCCTGGAGCATGTGAATGCCATCCAG 143  
 QY 36 GluAlaArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55  
 Db 144 GAGGCGCGCGTCTCTGAACCTGAGTGAAGACACTGCTGTGCTGAGATGAATGAACAGTA 203  
 QY 56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75  
 Db 204 GAAGTCATCTCAGAAATGTTTACCTCCAGGAGCGACCTGCTACAGACCGCCTGGAG 263  
 QY 76 LeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95  
 Db 264 CTGTACAAGCAGGCGCTCGGGGAGGCTCACCAAGCTCAAGGCGCCCTTGACCATGATG 323  
 QY 96 AlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIle 115  
 Db 324 GCCAGCCACTACAAGCAGCAGCTGCGCTCCAACCGCGAAATTCCTGTGCAACCCAGACT 383  
 QY 116 IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPheAsp 135  
 Db 384 ATCACCTTTGAAGATTTCAAGAGAACTTGAAGGACTTTCTGCTGTGTCATCCCTTTGAC 443  
 QY 136 CysTrpGluProValGlnGlu 142  
 Db 444 TGCTGGGAGCCAGTCCAGGAG 464

RESULT 9  
 CF370966  
 LOCUS  
 DEFINITION  
 TgESTyJ58e12.y1 Tg CAST Tachyzoite cDNA Library Toxoplasma gondii  
 cDNA clone TgESTyJ58e12.y1 5' similar to SW:CSF2 HUMAN P04141  
 GRANULOCYTE-MACROPHAGE COLONY-STIMULATING FACTOR PRECURSOR ; mRNA  
 sequence.  
 CF370966  
 CF370966.1 GI:34318212  
 EST.  
 Toxoplasma gondii  
 Toxoplasma gondii  
 Toxoplasma gondii  
 Eukaryota; Alveolata; Apicomplexa; Coccidia; Eimeriida;  
 Sarcocystidae; Toxoplasma.  
 1 (bases 1 to 592)  
 Tang,K., Cole,R., Fogarty,S., Sibley,L.D., Ajioka,J.A., White,M.,  
 Clifton,S., Page,D., Martin,J., Wyllie,T., Dante,M., Marra,M.,  
 Hillier,L., Kucaba,T., Theising,B., Bowers,Y., Gibbons,M.,  
 Ritter,E., Bennett,J., Franklin,C., Tsagarelisvili,R., Ronko,I.,  
 Kennedy,S., Maguire,L., Waterston,R. and Wilson,R.  
 Toxoplasma EST Project  
 Unpublished (2001)  
 Contact: Clifton, S.  
 Toxoplasma EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
 Tel: 314 286 1800

ACCESSION  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM

REFERENCE  
 AUTHORS

TITLE  
 JOURNAL  
 COMMENT



QY 75 luLeuTyLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetM 95  
 Db 272 AGCTGTACAAGCAGGGCTGGGGGCGACCTCACAGCTCAAGGGCCCTTGACCAIGA 331  
 QY 95 etAlaSerHisTyLysGlnHisCysProThrProGluThrSerCysAlaThrGlnI 115  
 Db 332 TGGCAGCCACTACAAGCAGCACTGCCCTCAACCCCGGAACTTCTGTGCAACCCAGA 391  
 QY 115 leIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPheA 135  
 Db 392 TTATCACCTTTGAAGTTTCAAGAGAACCTGAAGGACTTTCTGCTTGTATCCCTTTG 451  
 QY 135 spCysTrpGluProValGlnGlu 142  
 Db 452 ACTGCTGGGAGCCAGTCCAGGAG 474

## RESULT 11

CF370833 585 bp mRNA linear EST 27-AUG-2003  
 DEFINITION TgESTzyj55e12.y1 Tg CAST Tachyzoite cDNA Library Toxoplasma gondii  
 cDNA clone TgESTzyj55e12.y1 5' similar to SW-6292 HUMAN P04141  
 GRANULOCYTE-MACROPHAGE COLONY-STIMULATING FACTOR PRECURSOR ;, mRNA  
 sequence.

ACCESSION CF370833 GI:34318079  
 VERSION CF370833.1  
 KEYWORDS EST.  
 SOURCE Toxoplasma gondii

## ORGANISM

Toxoplasma gondii  
 Eukaryota; Alveolata; Apicomplexa; Coccidia; Eimeriida;

## REFERENCE

1 (bases 1 to 585)  
 Tang, K., Cole, R., Sibley, S., Fogarty, S., Ajioka, J. A., White, M.,  
 Clifton, S., Pape, D., Martin, J., Wylie, T., Dante, M., Marra, M.,  
 Hillier, L., Kucaba, T., Theising, B., Bowers, Y., Gibbons, M.,  
 Ritter, E., Bennett, J., Franklin, C., Tsagaris, V., Ronko, I.,  
 Kennedy, S., Maguire, L., Waterston, R. and Wilson, R.  
 Toxoplasma EST Project  
 Unpublished (2001)

## TITLE

## JOURNAL

## COMMENT

Toxoplasma EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: toxo@watson.wustl.edu

Contact David Sibley (toxost@borcim.wustl.edu) for further  
 information relating to organism, libraries, or clone availability.  
 Seq primer: -40UP from Gibco.

## FEATURES

## source

1. 585  
 /location/Qualifiers  
 /organism="Toxoplasma gondii"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:5811"  
 /clone="TgESTzyj55e12.y1"  
 /dev\_stage="Tachyzoite"

/lab\_host="Electron Blue cells (Stratagene)"

/clone\_lib="Tg CAST Tachyzoite cDNA Library"

/notes="Vector: Modified pBluescript (pBS SK+); Site 1:

BamHI; Site 2: EcoRI; The cDNA library was constructed by

Keliang Tang, and Robert Cole at Washington University.

cDNA was synthesized from poly(A)+ mRNA using the

template-switching PCR method (SMART cDNA Kit, BD

Biosciences). First strand cDNA was reverse transcribed

using the CDS III/3' primer and a 5' template switch

primer (Smart IV primer). The product of the first strand

synthesis was PCR amplified using the same primer set and

size selected, ligated into a modified pBluescript vector

(obtained from Michael White, Montana State University)

containing directional SfiI sites, and electroporated into

Electroten Blue cells. Vector: SfiI sites were added to

the multiple cloning region of pBluescript SK+ between the

BamHI/EcoRI sites. The modified polylinker has the  
 following sequence: 5'GAATTGGCCATTACGGCC(G)n-- insert--  
 GCGGCTCGCCCGACGGATCC3' where n=3-4 G nucleotides.  
 WARNING: the library contains a small percentage of cDNAs  
 derived from the human host cells. Library materials  
 provided by David Sibley, Washington University."

## ORIGIN

Alignment Scores:  
 Pred. No.: 4,01e-60 Length: 585  
 Score: 647.00 Matches: 123  
 Percent Similarity: 96.85% Conservative: 0  
 Best Local Similarity: 96.85% Mismatches: 4  
 Query Match: 84.58% Indels: 0  
 DB: 7 Gaps: 0

US-10-723-083-2 (1-142) x CF370833 (1-585)

QY 16 AlaProAlaArgSerProSerProSerThrGlnProTrpGluHisValAsnAlaIleGln 35  
 Db 86 GCACCCGCCCGCTGCCCGACGCCCGCCCGACGCGAGCCTGGGAGCATGTGAATGCCATCCAG 145  
 QY 36 GluAlaArgArgLeuLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55  
 Db 146 GAGGCCCGGCGTCTCTGAACCTGAGTAGAGACACTGCTGCTGAGATGAATTTAATAGTA 205  
 QY 56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75  
 Db 206 GAAGTCATCTCAGAAATGTTGACCTCCAGGAGCGACCTGCCTACAGACCCGCTGGAG 265  
 QY 76 LeuTyLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMet 95  
 Db 266 CTGTACAGCAGGGCTCGGGGCGAGCCCGCCAGCTCAAGGGCCCTTGACCAATGATG 325  
 QY 96 AlaSerHisTyLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIle 115  
 Db 326 GCCAGCCACTACAAGCAGCACTGCCCTCAACCCCGGAACTTCTGTGTGCAACCCAGACT 385  
 QY 116 IleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuLeuValIleProPheAsp 135  
 Db 386 ATCACCTTTGAAGTTTCAAGAGAACCTGAAGGACTTTCTGCTGTGCATCCCCCTTTCAC 445  
 QY 136 CysTrpGluProValGlnGlu 142  
 Db 446 TGCTGGGAGCCAGTCCAGGAG 466

## RESULT 12

## CD369973/c

## LOCUS

## DEFINITION

## ACCESSION

## CD369973

## KEYWORDS

## SOURCE

## ORGANISM

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## COMMENT

CD369973 718 bp mRNA linear EST 05-AUG-2004  
 UI-H-FT1-bke-o-08-0-UI-s1 NCI CGAP FT1 Homo sapiens cDNA clone  
 UI-H-FT1-bke-o-08-0-UI 3', mRNA sequence.

CD369973 GI:31154063

EST.

Homo sapiens (human)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 718)

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: [cgapb-remail.nih.gov](mailto:cgapb-remail.nih.gov)

Tissue Procurement: Dr. Gary W. Hunninghake, U of I

cDNA Library Preparation: Dr. M. Bento Soares, University of Iowa

DNA Sequencing by: Dr. M. Bento Soares, University of Iowa

Clone Distribution: Distribution information can be found at

<http://genome.uiowa.edu/distribution/cgap.html>

The following repetitive elements were found in this cDNA

sequence: 65-134, >(TAAA)n#Simple\_repeat

```

Seq primer: M13 FORWARD
POLYA=Yes.
FEATURES
    source
        Location/Qualifiers
            1..718
                /organism="Homo sapiens"
                /mol_type="mRNA"
                /db_xref="taxon:9606"
                /clone="UI-H-FTI-bke-o-08-0-UI"
                /tissue_type="Alveolar Macrophage"
                /dev_stage="Adult"
                /lab_host="DH10B (Life Technologies)"
                /clone_lib="NCI CGAP Ftl1"
                /notes="Organ: Lung; Vector: pT7T3-Pac (Pharmacia) with a
                modified polylinker; Site_1: EcoR I; Site_2: Not I;
                NCI CGAP_Ftl1 is a normalized cDNA library constructed from
                a pool of 81 RNA samples from Alveolar Macrophages
                challenged with different treatments. The mRNA samples
                were a mixture of these conditions (times refer to
                incubations following isolation by bronchoalveolar lavage)
                (some normal donor macrophages were cultured in some of
                the conditions, other donor macrophages in different
                conditions). The mRNA samples were pooled for library
                construction. Control 0 hours; LPS 100 ng/ml, 24 hours;
                PMA 10 ng/ml, 3 hours; PMA 10 ng/ml, 24 hours; Klebsiella
                moi 10, 3 hours; Klebsiella moi 10, 24 hours; Staph aureus
                vector (Ad5 CMV eGFP), moi 500, 3 hours; Adenoviral vector
                (Ad5 CMV eGFP), moi 500, 24 hours; wt adenovirus moi 500,
                3 hours; wt adenovirus moi 500, 24 hours; Ad vector + LPS
                3 hours; Ad vector + LPS 24 hours; wt adenovirus + LPS 3
                hours; wt adenovirus + LPS 24 hours. The library was
                normalized according to Bonaldo, Lennon and Soares, Genome
                Research, 6:791-806, 1996. First strand cDNA synthesis was
                primed with an oligo-dT primer containing a Not I site.
                Double stranded cDNA was ligated to an EcoR I adaptor,
                digested with Not I, and cloned directionally into
                pT7T3-Pac vector. The oligonucleotide used to prime the
                synthesis of first-strand cDNA contains a library tag
                sequence that is located between the Not I site and the
                (dT)18 tail. The sequence tag for this library is
                GGCCATGCCG. The tissue was provided by Dr. Gary W.
                Hunninghake of the University of Iowa.
                TAG_TISSUE=Human Lung Alveolar Macrophage
                TAG_LIB=UI-H-FTI
                TAG_SEQ=GGCCATGCCG"
    ORIGIN
        Alignment Scores:
        Pred. No.: 3,06e-59 Length: 718
        Score: 640.00 Matches: 125
        Percent Similarity: 99.21% Conservative: 0
        Best Local Similarity: 99.21% Mismatches: 1
        Query Match: 83.66% Indels: 1
        DB: 6 Gaps: 0

US-10-723-083-2 (1-142) x CD369973 (1-718)
QY 17 ProAlaArgSerProSerProThrGlnProTropGluHisValaenAlalleGlnGlu 36
Db 716 CCGCCGGCTCTGCAACCTGAGTACAGACACTGCTGCTGAGATGAATGCCATCCAGGAG 657
QY 37 AlaArgGluLeuLeuLeuLeuSerArgAspThrAlaAlaGluMetAsnGluThrValGlu 56
Db 656 GCCCGGGCTCTCTGCAACCTGAGTACAGACACTGCTGCTGAGATGAATGAACAGTAGAA 597
QY 57 ValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGluLeu 76
Db 596 GTCATCTCGAAATGTT-GACCTCCAGGAGCCGACCTGCTCAGACCCGCTGGAGCTG 538
QY 77 TyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMetAla 96
Db 537 TACAAGAGGGCTCGGGGCGAGCCCTCACCAGCTCAAGGGCCCTTGACCATGATGGCC 478

```

97 SerHisTyrIysGlnHisCysProProThrProGluThrSerCysAlaThrGlnIleIle 116  
 477 AGCCACTACAAGCAGCACTGCCCTCCAGCCCGAAACTTCTGTGCAACCCAGATTATC 418

117 ThrPheGluSerPheIysGluLeuLeuLeuAspPheLeuLeuValIleProPheAspCys 136  
 417 ACCTTTGAAGATTTCAAGAGAACCTTGAAGGACTTTTCTGCTTGTCTATCCCTTTGACTGC 359

137 TtpGluProValGlnGlu 142  
 357 TGGGAGCCAGTCCAGGAG 340

RESULT 13  
 BU633411/c  
 LOCUS  
 DEFINITION  
 UI-H-FL1-bgu-1-15-0-UI.s1 NCI CGAP FL1 Homo sapiens cDNA clone  
 UI-H-FL1-bgu-1-15-0-UI 3', mRNA sequence.  
 ACCESSION  
 BU633411 GI:23300666  
 VERSION  
 EST.  
 SOURCE  
 Homo sapiens (human)  
 ORGANISM  
 Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 1 (bases 1 to 701)  
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
 Tumor Gene Index  
 JOURNAL  
 Unpublished (1997)  
 COMMENT  
 Contact: Robert Strausberg, Ph.D.  
 Email: cgabbs@rmail.nih.gov  
 Tissue Procurement: James Martin  
 cDNA Library preparation: Dr. M. Bento Soares, University of Iowa  
 cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa  
 DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
 Clone Distribution: Clone distribution information can be obtained  
 from Dr. M. Bento Soares, bento-soares@uiowa.edu  
 The following repetitive elements were found in this cDNA  
 sequence: 1-46, >AT rich#Low\_complexity 60-129,  
 >(TAAA)n\$Simple repeat  
 Seq primer: M13\_FORWARD  
 POLYA=Yes.

FEATURES  
 source  
 1..701  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /clone="UI-H-FL1-bgu-1-15-0-UI"  
 /tissue\_type="Cell lines"  
 /dev\_stage="Adult"  
 /lab\_host="NCI CGAP FL1"  
 /clone\_lib="NCI CGAP FL1"  
 /notes="Organ: Chondrosarcoma; Vector: pT7T3-Pac  
 (Pharmacia) with a modified polylinker; Site\_1: EcoR I;  
 Site\_2: Not I; NCI CGAP FL1 is a normalized cDNA library  
 derived from a pool of mRNA obtained from 4 cell lines  
 from grade III chondrosarcoma tissues. The library was  
 constructed according to Bonaldo, Lennon and Soares,  
 Genome Research, 6:791-806, 1996. First strand cDNA  
 synthesis was primed with an oligo-dT primer containing a  
 Not I site. Double stranded cDNA was ligated to an EcoR I  
 adaptor, digested with Not I, and cloned directionally  
 into pT7T3-Pac vector. The oligonucleotide used to prime  
 the synthesis of first-strand cDNA contains a library tag  
 sequence that is located between the Not I site and the  
 (dT)18 tail. The sequence tag for this library is  
 GAGGTCGGTG. The cell lines were provided by Dr. James  
 Martin from the University of Iowa.  
 TAG\_TISSUE=Human Chondrosarcoma Grade 3 cell line mix  
 TAG\_LIB=UI-H-FL1  
 TAG\_SEQ=GAGTCCGTG"

ORIGIN

Alignment Scores:  
Pred. No.: 2,02e-56 Length: 701  
Score: 614.00 Matches: 120  
Percent Similarity: 98.36% Conservative: 0  
Best Local Similarity: 98.36% Mismatches: 2  
Query Match: 80.26% Indels: 1  
DB: 5 Gaps: 0

US-10-723-083-2 (1-142) x BU633411 (1-701)  
Qy 21 ProSerProSerThrGlnProTrpGluHisValAsnAlaIleGlnGluAlaArgArgLeu 40  
Db 699 CCCAGCCCCAGCAGCAGCGCTGGAGCATGTGAATCC-ATCCAGGAGGCGCGCTCTC 641  
Qy 41 LeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGluThrValGluValIleSerGlu 60  
Db 640 CTGAACCTGAGTGAAGACACTGCTGCTGAGATGAATGAACAGTAGAAGTCATCTCAGAA 581  
Qy 61 MetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGluLeuTyrLysGlnGly 80  
Db 580 ATGTTGACCTCCAGGAGCGACCTGCTACAGCCGCTGAGCTGTACAGAGGCG 521  
Qy 81 LeuArgGlySerLeuThrLysLeuLysGlyProLeuThrMetMetAlaSerHisTyrLys 100  
Db 520 CTCGGGGCAGCCTCACCAAGCTCAAGGCGCCCTTGACCATGATGCCAGCCACTACAAG 461  
Qy 101 GlnHisCysProProThrProGluThrSerCysAlaThrGlnIleIleThrPheGluSer 120  
Db 460 CAGCAGTGCCCTCAACCCCGGAACCTCTCTGTGCAACCCAGATATACCTTTGAAGT 401  
Qy 121 PheLysGluAsnLeuLysAspPheLeuValIleProPheAspCysTrpGluProVal 140  
Db 400 TTCAAGAGAACCTGAAGGACTTCTCTGTGTCATCCCTTTGACTGCTGGGAGCCAGTC 341  
Qy 141 GlnGlu 142  
Db 340 CAGGAG 335

RESULT 14  
CA307828/c  
LOCUS CA307828 666 bp mRNA linear EST 05-AUG-2004  
DEFINITION UI-H-FTI-bhx-f-10-0-UI.s1 NCI\_CGAP\_FTI Homo sapiens cDNA clone  
CA307828  
ACCESSION CA307828.1 GI:24470882  
VERSION  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 666)  
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),  
Tumor Gene Index  
Unpublished (1997)  
Contact: Robert Strausberg, Ph.D.  
Email: [cgaps@mail.nih.gov](mailto:cgaps@mail.nih.gov)  
Tissue Procurement: Dr. Gary W. Hunninghake, U of I  
CDNA Library Preparation: Dr. M. Bento Soares, University of Iowa  
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
Clone Distribution: Clone distribution information can be obtained  
from Dr. M. Bento Soares, [bento-soares@uiowa.edu](mailto:bento-soares@uiowa.edu)  
The following repetitive elements were found in this cDNA  
sequence: i-50, >AT\_richLow\_complexity 64-133,  
>(TAAA)n#Simple\_repeat  
Seq primer: M13 FORWARD  
POLYA=Yes.

Location/Qualifiers  
1. .666  
/organism="Homo sapiens"  
/mol\_type="mRNA"

FEATURES  
source

ORIGIN  
Alignment Scores:  
Pred. No.: 6.88e-51 Length: 666  
Score: 563.00 Matches: 108  
Percent Similarity: 99.08% Conservative: 0  
Best Local Similarity: 99.08% Mismatches: 1  
Query Match: 73.59% Indels: 0  
DB: 6 Gaps: 0

US-10-723-083-2 (1-142) x CA307828 (1-666)  
Qy 34 IleGlnGluAlaArgArgLeuAsnLeuSerArgAspThrAlaAlaGluMetAsnGlu 53  
Db 665 ATCCAGGAGGCGCGCTCTCTGAACCTGAGTAGAGACACTGCTGTGAGATGAATGAA 606  
Qy 54 ThrValGluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArg 73  
Db 605 ACAGTAGAAGTCATCTCAGAAATGTTTACCTCCAGGAGCGCCACTGCTACAGACCCGC 546  
Qy 74 LeuGluLeuTyrLysGlnGlyLeuArgGlySerLeuThrLysLeuLysGlyProLeuThr 93  
Db 545 TTGGAGCTGTACAAGCAGGCGCTGCGGGGCGACCTCACCAAGCTCAAGGGCCCTTGACC 486  
Qy 94 MetMetAlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThr 113  
Db 485 ATGATGGCCAGCCANTCAAGCAGCAGCTCCCTCCACCCCGGAACTTCTCTGTGCAACC 426  
Qy 114 GlnIleIleThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIlePro 133  
Db 425 CAGATTATCACCTTTGAAGATTCAAGAGAACCTGAGGAGACTTTCTGCTTGTCTATCCC 366  
Qy 134 PheAspCysTrpGluProValGlnGlu 142

/db\_xref="taxon:9606"  
/clone="UI-H-FTI-bhx-f-10-0-UI"  
/tissue\_type="Alveolar Macrophage"  
/dev\_stage="Adult"  
/lab\_host="DH10B (Life Technologies)"  
/clone\_lib="NCI CGAP\_FTI"  
/note="Organ: Lung; Vector: pT7T3-Pac (Pharmacia) with a modified polylinker; Site 1: EcoR I; Site 2: Not I; NCI\_CGAP\_FTI is a normalized cDNA library constructed from a pool of 81 RNA samples from Alveolar Macrophages challenged with different treatments. The mRNA samples were a mixture of these conditions (times refer to incubations following isolation by bronchoalveolar lavage) (some normal donor macrophages were cultured in some of the conditions, other donor macrophages in different conditions). The mRNA samples were pooled for library construction. Control 0 hours; control 3 hours; control 24 hours; LPS 100 ng/ml, 3 hours; LPS 100 ng/ml, 24 hours; PMA 10 ng/ml, 3 hours; PMA 10 ng/ml, 24 hours; Klebsiella moi 10, 3 hours; Klebsiella moi 10, 24 hours; Adenovirus moi 10, 3 hours; Staph aureus moi 10, 24 hours; Adenoviral vector (Ad5 CMV eGFP), moi 500, 3 hours; Adenoviral vector (Ad5 CMV eGFP), moi 500, 24 hours; wt adenovirus moi 500, 3 hours; wt adenovirus moi 500, 24 hours; Ad vector + LPS 3 hours; Ad vector + LPS 24 hours; wt adenovirus + LPS 3 hours; wt adenovirus + LPS 24 hours. The library was normalized according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pT7T3-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (GT)18 tail. The sequence tag for this library is GCCCATGCCG. The tissue was provided by Dr. Gary W. Hunninghake of the University of Iowa.  
TAG\_LIB=UI-H-FTI  
TAG\_SEQ=GGCATGCCG"

```

Db      365 TTTGACTGCTGGGAGCAGTCCAGGAG 339
|||||
RESULT 15
AA995402
LOCUS   475 bp mRNA linear EST 27-AUG-1998
DEFINITION
or74f05.s1 NCI CGAP Lu5 Homo sapiens cDNA clone IMAGE:1601601 3'
similar to gb:M11220 GRANULOCYTE-MACROPHAGE COLONY-STIMULATING
FACTOR PRECURSOR (HUMAN); mRNA sequence.
ACCESSION
AA995402
VERSION
AA995402.1 GI:3181891
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 475)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
AUTHORS
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished (1997)
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgabs-r@mail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
DNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www.bio.llnl.gov/bbrp/image/image.html
Insert length: 747 Std Error: 0.00
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 308.

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## FEATURES

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Location/Qualifiers
1..475
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1601601"
/tissue_type="carcinoid"
/lab_host="DH10B"
/clone_lib="NCI CGAP Lu5"
/note="Organ: lung; Vector: pT73D-Pac (Pharmacia) with a
modified polylinker; 1st strand cDNA was prepared from
neuroendocrine lung carcinoid, and was then primed with a
Not I - oligo(dT) primer. Double-stranded cDNA was ligated
to Eco RI adaptors (Pharmacia), digested with Not I and
cloned into the Not I and Eco RI sites of the modified
pT73 vector. Library is normalized. Library was
constructed by Bento Soares and M. Fatima Bonaldo."

```

## ORIGIN

```

Alignment Scores:
Pred. No.: 12e-50 Length: 475
Score: 559.00 Matches: 119
Percent Similarity: 94.49% Conservative: 1
Best Local Similarity: 93.70% Mismatches: 6
Query Match: 73.07% Indels: 3
DB: 1 Gaps: 0

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US-10-723-083-2 (1-142) x AA995402 (1-475)

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QY      16 AlaProAlaArgSerProSerThrGlnProTrpGluHisValaenAlaIleGln 35
|||||
Db      55 GCACCCGCCCGCTCGCCAGCCAGCAGCCCTGGGAGCATGTGATGC-ATCCAG 113
|||||
QY      36 GluAlaArgLeuLeuLeuLeuLeuSerArgAspThrAlaAlaGluMetAsnGluThrVal 55
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Db      114 GAGGCCCGCGCTCTCTGAACCTGAGTAGAGACACTGCTGAGATGAATGAACAGTA 173
|||||
QY      56 GluValIleSerGluMetPheAspLeuGlnGluProThrCysLeuGlnThrArgLeuGlu 75
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Search completed: March 11, 2005, 20:24:33  
Job time : 2828 secs

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Db      174 GAAGTCATCTCAGAAATGTTTGACCTCCAGGAGCCGACCTGCCTACAGACCCGCTGGAG 233
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QY      76 LeuTyrLysGlnGlyLeuArgGlySer-LeuThrLysLeuLysGlyProLeuThrMetMe 95
|||||
Db      234 CTGTACAAGCAGGGCCTGGGTGGCAGTCTCTACCAAGCTCAGGGGCCCTTTGACCATGAT 293
|||||
QY      95 tAlaSerHisTyrLysGlnHisCysProProThrProGluThrSerCysAlaThrGlnI 115
|||||
Db      294 GGCACGCCACTACAGCAGCAGCTGCCCTCCAAACCCGGAACCTTCCTTTTGC AAC-CAGAT 352
|||||
QY      115 eileThrPheGluSerPheLysGluAsnLeuLysAspPheLeuValIleProPheAs 135
|||||
Db      353 TATCACCTCTTGNAGTTTCAAGAGAACCTGAAGGACTTTCTGCTGTGTCATCCACTTTGA 412
|||||
QY      135 pCysTrpGluProValGln 141
|||||
Db      413 CTGCTGGGAGCCAGTCAGG 431
|||||

```

